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(54) Title: RECOMBINANT HK2 POLYPEPTIDE			
<p style="text-align: right;">41</p> <p>hK2: <u>IVGGWECEKHSQFPQVAVWSHGWAHCGGVLVHPQWVLTAAHCLKKN SQVWLGRHN</u></p> <p>hK3: <u>#####L A R R V #####IRNK VIL #####S</u></p> <p style="text-align: right;">56</p> <p><u>LFEPEDTGQRVPVSHSFPHPLYNMSLLKHQSLRPDEDSSHDLM LRLSEPARIT</u></p> <p><u>#####V F Q #####D #####N R F #####G D #####E L</u></p> <p style="text-align: right;">110</p> <p><u>DVVKVLGLPTQEPALGTTTCYASGWGSI EPEFLRPRSLQCVSLHLLSNDHCA</u></p> <p><u>#####M D #####T P K K #####Q #####V I #####V</u></p> <p style="text-align: right;">153</p> <p style="text-align: right;">162 167</p> <p><u>RAYSEKVTEFMLCAGLWTGGKDTCCGDSGGPLVCNGVLQGITSWGPEPCALPEKP</u></p> <p><u>OVHPO #####X #####R #####S #####S #####R</u></p> <p style="text-align: right;">217 237</p> <p><u>AVYTKVVHYRKWIKDTIAANP</u></p> <p><u>SL #####V</u></p>			
(57) Abstract			
<p>An isolated, substantially homogenous hK2 polypeptide is provided as well as isolated nucleic acid molecules encoding hK2 polypeptide, including (a) a cDNA molecule comprising the nucleotide sequence of the coding region of human hK2 gene; (b) a DNA molecule capable of hybridizing under stringent conditions to a molecule of (a); and (c) a genetic variant of any of the DNA molecules of (a) and (b) which encodes of polypeptide processing an antigenic function of naturally occurring polypeptide.</p>			

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RECOMBINANT HK2 POLYPEPTIDE

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Background of the Invention

The glandular kallikreins are a subgroup of serine proteases which are involved in the post-translational processing of specific polypeptide precursors to their biologically active forms. The human kallikrein gene family consists of three members: prostate-specific antigen, human glandular kallikrein, and pancreatic/renal kallikrein. See J.A. Clements, Endocr. Rev., 10, 393 (1989) and T.M. Chu et al. (U.S. Patent No. 4,446,122). A common nomenclature for these members of the tissue (glandular) kallikrein gene families was recently adopted by T. Berg et al., in Recent Progress on Kinins: Biochemistry and Molecular Biology of the Kallikrein-Kinin System. Agents and Actions Supplements, Vol. I., H. Fritz et al., eds., Birkhauser Verlag, Basel (1992), and is defined in Table I, below.

TABLE 1

20

The Human Tissue Kallikrein Gene Family
(approved species designation: HSA)

25	New Designation	Previous Designations	mRNA/cDNA	Protein	New Protein Designation
30	hKLK1	KLK1 hRKALL	λ HK1 and phKK25 cDNAs	tissue kallikrein (renal/ pancrease/salivary)	hK1
35	hKLK2	KLK2 hGK-1 hKK-3		prostate-specific glandular kallikrein	hK2
40	hKLK3	PSA PA APS	λ HPSA-1 and PSA cDNAs	PSA (prostate-specific antigen)	hK3

The DNA sequence homology between hCLK2 and hCLK3 (exon regions) is 80%, whereas the homology between hCLK2 and hCLK1 is 65%. The deduced amino acid sequence homology of hK2 to hK1 is 57%. Amino acid sequences deduced by L.J. Schedlich et al., *DNA*, 6, 429 (1987) and B.J. Morris, *Clin. Exp. Pharmacol. Physiol.* 16, 345 (1989) indicate that hK2 may be a trypsin-like serine protease, whereas hK3 (PSA) is a chymotrypsin-like serine protease. Therefore, if hK2 is indeed secretory, it may have a different physiological function than hK3.

The hCLK2 gene is located about 12 kbp downstream from the hCLK3 gene in a head-to-tail fashion on chromosome 19. (P.H. Riegman et al., *FEBS Lett.* 247, 123, (1989)). The similarities of gene structure and deduced amino acid sequences of these human kallikreins suggest that their evolution may involve the same ancestral gene. Most interestingly, as reported by Morris, cited *supra*; P. Chapdelaine, *FEBS Lett.* 236, 205 (1988); and Young, *Biochemistry*, 31, 1952 (1992), both hK2 and hK3 may be expressed only in the human prostate, while expression of hK1 is limited to the pancreas, submandibular gland, kidney, and other nonprostate tissues.

Tremendous interest has been generated in hK3 (PSA) because of the important role it plays as a marker to detect and to monitor progression of prostate carcinoma. Its usefulness as a marker is based on the elevated serum concentration of circulating hK3 proteins which are frequently associated with prostatic cancer. The serum concentration of hK3 has been found to be proportional to the cancer mass in untreated patients, but is also proportional to the volume of hyperplastic tissue in patients with benign prostatic hyperplasia (BPH). The serum levels of hK3 become reduced following prostate cancer therapy.

Despite the information which can be ascertained about hK2 from the genomic DNA sequence, very little is known about the hK2 polypeptide itself. The reason for this is that the protein has not been purified and characterized. Thus, a need exists for a method to obtain hK2 polypeptide and related polypeptides in sufficient quantity and purity for characterization and for use as therapeutic/diagnostic agents or reagents.

Summary of the Invention

The present invention provides an isolated, substantially homogenous hK2 polypeptide. As used herein, in the term "hK2 polypeptide" includes pre-pro hK2, pro hK2 and mature hK2 polypeptides. Pre-pro hK2 is secreted by the cell *in vivo*, and is cleaved during secretion to yield pro hK2, which is then enzymatically cleaved in the extracellular environment to yield "mature" hK2. Most preferably, the hK2 polypeptide is contiguous in amino acid sequence with SEQ ID NO: 16, SEQ ID NO: 6 or SEQ ID NO: 10.

The present invention also provides isolated nucleic acid molecules encoding hK2 polypeptide, including (a) a cDNA molecule comprising the nucleotide sequence of the coding region of the hK2 gene; (b) a DNA molecule capable of hybridizing under stringent conditions to a nucleotide sequence complementary to the nucleotide sequence of (a); and (c) a genetic variant of any of the DNA molecules of (a) and (b) which encodes of polypeptide processing an antigenic function of naturally occurring hK2 polypeptide. Preferably, the nucleic acid comprises a discrete, isolated DNA or RNA molecule encoding the complete hK2 polypeptide, which can include the pre-pro, pro or mature forms. Most preferably, the nucleic acid is a DNA sequence contiguous with SEQ ID NO: 5, 7 or 9, i.e., as shown in Figs. 5, 6 or 7. These DNA sequences can be produced using the polymerase chain reaction (PCR), and novel oligonucleotide primers employed in the synthesis are also an embodiment of the invention.

The nucleic acid sequence also can comprise a promoter operably linked to the nucleic acid sequence. Therefore, the invention also comprises a chimeric expression vector comprising the above-described nucleic acid sequence, operationally linked to control sequences recognized by a host cell transformed with the vector, as well as said transformed host cell, and methods of its preparation and use to produce recombinant hK2. Thus, the present invention also provides a method of using a nucleic acid molecule, such as a cDNA clone encoding hK2 polypeptide, comprising expressing the nucleic acid molecule in a cultured host cell transformed, preferably stably transformed, with a chimeric expression vector comprising said nucleic acid

molecule operably linked to control sequences recognized by the host cell transformed with the vector; and recovering the hK2 polypeptide from the transgenic host cell, i.e., from the culture medium. As used herein, the term "chimeric" means that the vector comprises DNA from at least two different
5 species, or comprises DNA from the same species, which is linked or associated in a manner which does not occur in the "native" form of said species.

More specifically, E. coli and baculovirus insect cells systems have been employed to produce hK2 polypeptides in two forms, i.e. pre-pro
10 hK2 (pphK2) and mature hK2 (mhK2). Thus, the present invention provides the first example of the overexpression of hK2 in heterologous systems. However, although pphK2 produced in E. coli has proven to be an invaluable resource for generating antibodies to the denatured form of the protein, it is desirable to both discern the steps involved in the biosynthesis of hK2 and to
15 obtain antibodies specific for the fully processed and secreted form of the protein. Therefore, mammalian cell systems have been employed to produce hK2 polypeptides. Thus, the present invention also provides the first example of the expression of hK2 in mammalian cells and purification and characterization of the secreted protein.

20 The high degree of amino acid sequence homology of hK2 with hK3 indicates that measuring serum concentrations of both proteins may be useful in the diagnosis and monitoring of prostate cancer. For example, the antibodies developed against hK3 now used in these assays could theoretically also recognize hK2, because of mutual contamination in the antigenic
25 preparations used to develop the anti-hK3 antibodies or because of antibody cross-reactivity between these two proteins. This could account for the substantial percentage of false positive results which are observed in current hK3 assays. On the other hand, if circulating hK2 levels are also elevated above baseline levels in prostate cancer patients, detection of hK2 by hK2-
30 specific antibodies could provide an alternative, confirmatory assay for prostate cancer.

Therefore, hK2 polypeptide, as well as variants and subunits thereof, produced by the present method can be used to produce populations of antibodies that, in turn, can be used as the basis for assays to detect and quantify hK2 polypeptide (or "protein") in samples derived from tissues such as prostate carcinomas, cells such as prostate cell lines, or from fluids such as seminal fluid or blood. Thus, the present invention also provides populations of monoclonal or polyclonal antibodies that specifically bind to hK2 polypeptide, while not significantly binding to hK3. The term "significantly" is defined by reference to the comparative assays discussed below. These antibodies can also be used in affinity chromatography, to purify mammalian hK2 from natural sources. The isolated, substantially homogeneous hK2 can also be employed as a component in diagnostic assays for "native" hK2 in samples derived from human tissues or physiological fluids. For example, the recombinant hK2 can be bound to a detectable label and employed in competitive immunoassays for hK2, as described in U.S. patent application Serial No. 08/096,946, filed July 22, 1993.

As used herein with respect to the present invention, the terms "hK2 polypeptide," "hK2 protein," and "hK2" are considered to refer to identical human materials, unless otherwise indicated.

Brief Description of the Figures

Figure 1 depicts a time course study of recombinant pphK2 in Sf9 cells infected with recombinant pphK2 virus. At each of the time points cells were depleted of methionine and cysteine for 1 hour in deficient media and then supplemented with [³⁵S]-methionine and [³⁵S]-cysteine. Protein was determined by Bradford assay. Aliquots of protein (20 µg) were loaded onto a 12% Tris-Glycine SDS gel. A Phosphorimager cassette was exposed overnight. The band of interest is indicated with an arrow. w.t.: wild type.

Figure 2 depicts the detection of recombinant mhK2 in cell lysate fractions. Sf9 cells were infected either with recombinant mhK2, wild type or left uninfected for 48 hours. Methionine and cysteine pools were depleted for 1 hour in deficient media. Cells were supplemented with [³⁵S]-

methionine and [³⁵S]-cysteine for 6 hours. Cells were separated into soluble and insoluble fractions using H₂O and repeated freeze/thaw conditions. Aliquots of protein (50 µg per lane) were loaded onto a 10% Tris-Glycine SDS gel and electrophoresed. The gel was dried and exposed to x-ray film
5 for 2 days. The band of interest is indicated with an arrow.

Figure 3 depicts the expression of recombinant pphK2 in *E. coli*. *E. coli* strain BL21 (DE3) LysS harboring pBppHK2 was grown in LB media to O.D.₆₀₀ 0.2 and incubated without (lane 2, not-induced (N)) or with (lane 3, induced (I)) 0.4 mM IPTG for 2 hrs. Cells were lysed in sample
10 buffer and subjected to SDS/PAGE on a 4-20% gradient gel. Protein bands were visualized by staining the gel with Coomassie blue.

Figure 4 depicts the amino acid sequences of mature hK2 (deduced from cDNA sequence, SEQ ID NO: 16) and hK3 (SEQ ID NO: 1). Underlined sequences denote nonhomologous regions that can be used for
15 preparation of antibodies specific to hK2.

Figure 5 depicts pphK2 cDNA containing a BamHI site at the 5' end and a PstI site at the 3' end (SEQ ID NO: 5) (coding strand is numbered) as well as the amino acid sequence of pre-pro hK2 encoded thereby (SEQ ID NO: 6). The amino acid sequences of pro hK2 and mature
20 hK2 are also shown on the Figure.

Figure 6 depicts mhK2 cDNA containing an EcoRI site at the 5' end and PstI site at the 3' end (SEQ ID NO: 7), as well as the corresponding amino acid sequence (SEQ ID NO: 8) which encompasses the amino acid sequence of mhK2 polypeptide.

25 Figure 7 depicts pro hK2 DNA (SEQ ID NO: 9) (coding strand is numbered) and the amino acid sequence of pro hK2 (SEQ ID NO: 10).

Figure 8 depicts a gel confirming the expression of recombinant pphK2 in a mammalian cell line. AV12-pGThK2 (Lane 4-6) and AV12-pGT-d (Lane 3) clonal cell lines were grown in D10F media. About 300µl of
30 spent medium from the above clones were concentrated and subjected to SDS/PAGE along with See Blue MW marker (lane 1) and pphK2 lysate from *E. coli* cells (lane 2). The gel was blotted onto nitrocellulose paper and

immunoblotted using a 1/1000 dilution of anti-phK2 rabbit antiserum. HRP-goat anti-rabbit was used as the secondary probe and the blot was developed by DAB plus H₂O₂. Lane 3 (AV12-pGT-d) is AV12 transfected with vector without insert.

5 Figure 9 depicts the DEAE chromatography of AV12 media. The sample was applied in a bicarbonate buffer, pH 8 and eluted with a salt gradient. The solid line is the A₂₈₀ elution profile. The triangle line represents the ELISA assay of individual samples which had been dried onto microtiter plates and developed with rabbit anti-hK2 antibody.

10 Figure 10 depicts the hydrophobic interaction profile of DEAE fractions. The fractions were pooled, concentrated and applied to an HIC column in 1.2 M sodium sulfate, and eluted with a decreasing salt gradient. The solid line is A₂₈₀ and the triangle line shows the ELISA assay profile of the fractions using rabbit anti-hK2 antibody.

15 Figure 11 depicts the Size Exclusion Chromatography of HIC purified prohK2, in particular, the A₂₈₀ profile of 22 min peak eluted off HIC column. The 19.4 min peak appears homogeneous by SDS-PAGE. After this peak was lyophilized, the N-terminal sequence and amino acid composition confirmed its identity as the pro form of hK2.

20 Figure 12 depicts the SDS/PAGE analysis of prohK2 and PSA. 1.5µg of purified phK2 or PSA was boiled in sample buffer containing (R) or not containing (N) 1% BME. Samples were subjected to SDS/PAGE on a 4-20% gel. The protein bands were visualized by staining the gel with silver.

25 Detailed Description of the Invention

As used herein, the term "hK2 polypeptide" preferably encompasses the recombinant pre-pro, pro and mature hK2 polypeptides. As proposed herein, a mature hK2 polypeptide having the amino acid sequence shown in Fig. 4 (SEQ ID NO: 16), as well as "variant" polypeptides which
30 share at least 90% homology with SEQ ID NO: 16 in the regions which are substantially homologous with hK3, i.e., which regions are not identified by bars as shown in Fig. 4. Such hK2 polypeptides also possess antigenic

function in common with the mature hK2 molecule of Fig. 4, in that said polypeptides are also definable by antibodies which bind specifically thereto, but which do not cross-react with hK3 (or hK1). Preferably, said antibodies react with antigenic sites or epitopes that are also present on the mature hK2 molecule of Fig. 4. Antibodies useful to define common antigenic function are described in detail in Ser. No. 08/096,946, i.e., polyclonal antisera prepared in vivo against hK2 submit 41-56.

"Isolated hK2 nucleic acid" is RNA or DNA containing greater than 15, preferably 20 or more, sequential nucleotide bases that encode a biologically active hK2 polypeptide or a variant fragment thereof, that is complementary to the non-coding strand of the native hK2 polypeptide RNA or DNA, or hybridizes to said RNA or DNA and remains stably bound under stringent conditions. Thus, the RNA or DNA is isolated in that it is free from at least one contaminating nucleic acid with which it is normally associated in the natural source and is preferably substantially free of any other mammalian RNA or DNA. The phrase "free from at least one contaminating source nucleic acid with which it is normally associated" includes the case where the nucleic acid is reintroduced into the source or natural cell but is in a different chromosomal location or is otherwise flanked by nucleic acid sequences not normally found in the source cell. An example of isolated hK2 nucleic acid is RNA or DNA that encodes a biologically active hK2 polypeptide sharing at least 90% sequence identity with the hK3-homologous regions of the hK2 peptide of Fig. 4, as described above. The term "isolated, substantially homogenous" as used with respect to an hK2 polypeptide is defined in terms of the methodologies discussed herein below.

As used herein, the term "recombinant nucleic acid," i.e., "recombinant DNA" refers to a nucleic acid, i.e., to DNA that has been derived or isolated from any appropriate tissue source, that may be subsequently chemically altered *in vitro*, and later introduced into target host cells, such as cells derived from animal, plant, insect, yeast, fungal or bacterial sources. An example of recombinant DNA "derived" from a source, would be a DNA sequence that is identified as a useful fragment encoding

hK2, or a fragment or variant thereof, and which is then chemically synthesized in essentially pure form. An example of such DNA "isolated" from a source would be a useful DNA sequence that is excised or removed from said source by chemical means, e.g. by the use of restriction
5 endonucleases, so that it can be further manipulated, e.g., amplified, for use in the invention, by the methodology of genetic engineering.

Therefore, "recombinant DNA" includes completely synthetic DNA sequences, semi-synthetic DNA sequences, semi-synthetic DNA sequences, DNA sequences isolated from biological sources, and DNA
10 sequences derived from introduced RNA, as well as mixtures thereof. Generally, the recombinant DNA sequence is not originally resident in the genome of the host target cell which is the recipient of the DNA, or it is resident in the genome but is not expressed.

The recombinant DNA sequence, used for transformation
15 herein, may be circular or linear, double-stranded or single-stranded. Generally, the DNA sequence is in the form of chimeric DNA, such as plasmid DNA, that can also contain coding regions flanked by control sequences which promote the expression of the recombinant DNA present in the resultant cell line. For example, the recombinant DNA may itself
20 comprise a promoter that is active in mammalian cells, or may utilize a promoter already present in the genome that is the transformation target. Such promoters include the CMV promoter, as well as the SV 40 late promoter and retroviral LTRs (long terminal repeat elements). Aside from recombinant DNA sequences that serve as transcription units for hK2 or
25 portions thereof, a portion of the recombinant DNA may be untranscribed, serving a regulatory or a structural function.

"Control sequences" is defined to mean DNA sequences necessary for the expression of an operably linked coding sequence in a particular host organism. The control sequences that are suitable for
30 prokaryotic cells, for example, include a promoter, and optionally an operator sequence, and a ribosome binding site. Eukaryotic cells are known to utilize promoters, polyadenylation signals, and enhancers.

"Operably linked" is defined to mean that the nucleic acids are placed in a functional relationship with another nucleic acid sequence. For example, DNA for a presequence or secretory leader is operably linked to DNA for a polypeptide if it is expressed as a preprotein that participates in the secretion of the polypeptide; a promoter or enhancer is operably linked to a coding sequence if it affects the transcription of the sequence; or a ribosome binding site is operably linked to a coding sequence if it is positioned so as to facilitate translation. Generally, "operably linked" means that the DNA sequences being linked are contiguous and, in the case of a secretory leader, contiguous and in reading phase. However, enhancers do not have to be contiguous. Linking is accomplished by ligation at convenient restriction sites. If such sites do not exist, the synthetic oligonucleotide adaptors or linkers are used in accord with conventional practice.

Aside from recombinant DNA sequences that serve as transcription units for hK2 or portions thereof, a portion of the recombinant DNA may be untranscribed, serving a regulatory or a structural function.

The recombinant DNA to be introduced into the cells further will generally contain either a selectable marker gene or a reporter gene or both to facilitate identification and selection of transformed cells from the population of cells sought to be transformed. Alternatively, the selectable marker may be carried on a separate piece of DNA and used in a co-transformation procedure. Both selectable markers and reporter genes may be flanked with appropriate regulatory sequences to enable expression in the host cells. Useful selectable markers are well known in the art and include, for example, antibiotic and herbicide-resistance genes, such as neo, hpt, dhfr, bar, aroA, dapA and the like.

Reporter genes are used for identifying potentially transformed cells and for evaluating the functionality of regulatory sequences. Reporter genes which encode for easily assayable proteins are well known in the art. In general, a reporter gene is a gene which is not present in or expressed by the recipient organism or tissue and which encodes a protein whose expression is manifested by some easily detectable property, e.g., enzymatic activity.

Preferred genes include the chloramphenicol acetyl transferase gene (cat) from Tn9 of *E. coli*, the beta-glucuronidase gene (gus) of the *uidA* locus of *E. coli*, and the luciferase gene from firefly *Photinus pyralis*. Expression of the reporter gene is assayed at a suitable time after the DNA has been introduced
5 into the recipient cells.

Other elements functional in the host cells, such as introns, enhancers, polyadenylation sequences and the like, may also be a part of the recombinant DNA. Such elements may or may not be necessary for the function of the DNA, but may provide improved expression of the DNA by
10 affecting transcription, stability of the mRNA, or the like. Such elements may be included in the DNA as desired to obtain the optimal performance of the transforming DNA in the cell.

The general methods for constructing recombinant DNA which can transform target cells are well known to those skilled in the art, and the
15 same compositions and methods of construction may be utilized to produce the DNA useful herein. For example, J. Sambrook et al., Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Laboratory Press (2d ed., 1989), provides suitable methods of construction.

The recombinant DNA can be readily introduced into the target
20 cells by transfection with an expression vector comprising cDNA encoding hK2, for example, by the modified calcium phosphate precipitation procedure of C. Chen et al., Mol. Cell. Biol., 7, 2745 (1987). Transfection can also be accomplished by lipofectin, using commercially available kits, e.g., provided by BRL.

Suitable host cells for the expression of hK2 polypeptide are
25 derived from multicellular organisms. Such host cells are capable of complex processing and glycosylation activities. In principle, any higher eukaryotic cell culture is workable, whether from vertebrate or invertebrate culture. Examples of invertebrate cells include plant and insect cells. Numerous
30 baculoviral strains and variants and corresponding permissive insect host cells from hosts such as *Spodoptera frugiperda* (caterpillar), *Aedes aegypti* (mosquito), *Aedes albopictus* (mosquito), *Drosophila melanogaster* (fruitfly),

and *Bombyx mori* have been identified. See, e.g., Luckow et al., Bio/Technology, 6; 47 (1988); Miller et al., in Genetic Engineering, J. K. Setlow et al., eds., Vol. 8 (Plenum Publishing, 1986), pp. 277-279; and Maeda et al., Nature, 315:592 (1985). A variety of viral strains for transfection are
5 publicly available, e.g., the L-1 variant of *Autographa californica* NPV and the Bm-5 strain of *Bombyx mori* NPV, and such viruses may be used, preferably for transfection of *Spodoptera frugiperda* cells.

Recovery or isolation of a given fragment of DNA from a restriction digest can employ separation of the digest on polyacrylamide or
10 agarose gel by electrophoresis, identification of the fragment of interest by comparison of its mobility versus that of marker DNA fragments of known molecular weight, removal of the gel section containing the desired fragment, and separation of the gel from DNA. For example, see Lawn et al., Nucleic
Acids Res., 9, 6103-6114 (1981), and Goeddel et al., Nucleic Acids Res., 8,
15 4057 (1980).

"Southern analysis" or "Southern blotting" is a method by which the presence of DNA sequences in a restriction endonuclease digest of DNA or DNA-containing composition is confirmed by hybridization to a known, labeled oligonucleotide or DNA fragment. Southern analysis typically
20 involves electrophoretic separation of DNA digests on agarose gels, denaturation of the DNA after electrophoretic separation, and transfer of the DNA to nitrocellulose, nylon, or another suitable membrane support for analysis with a radiolabeled, biotinylated, or enzyme-labeled probe as described in sections 9.37-9.52 of Sambrook et al., *supra*.

25 "Northern analysis" or "Northern blotting" is a method used to identify RNA sequences that hybridize to a known probe such as an oligonucleotide, DNA fragment, cDNA or fragment thereof, or RNA fragment. The probe is labeled with a radioisotope such as ³²P, by biotinylation or with an enzyme. The RNA to be analyzed can be usually electrophoretically
30 separated on an agarose or polyacrylamide gel, transferred to nitrocellulose, nylon, or other suitable membrane, and hybridized with the probe, using

standard techniques well known in the art such as those described in sections 7.39-7.52 of Sambrook et al., *supra*

"Polymerase chain reaction" or "PCR" refers to a procedure or technique in which amounts of a preselected piece of nucleic acid, RNA and/or DNA, are amplified as described in U.S. Patent No. 4,683,195. Generally, sequence information from the ends of the region of interest or beyond is employed to design oligonucleotide primers. These primers will be identical or similar in sequence to opposite strands of the template to be amplified. PCR can be used to amplify specific RNA sequences, specific DNA sequences from total genomic DNA, and cDNA transcribed from total cellular RNA, bacteriophage or plasmid sequences, and the like. See generally Mullis et al., *Cold Spring Harbor Symp. Quant. Biol.*, **51**, 263 (1987); Erlich, ed., *PCR Technology*, (Stockton Press, NY, 1989).

"Stringent conditions" are those that (1) employ low ionic strength and high temperature for washing, for example, 0.015 M NaCl/0.0015 M sodium citrate (SSC); 0.1% sodium lauryl sulfate (SDS) at 50°C, or (2) employ during hybridization a denaturing agent such as formamide, for example, 50% (vol/vol) formamide with 0.1% bovine serum albumin/0.1% Ficoll/0.1% polyvinylpyrrolidone/50 mM sodium phosphate buffer at pH 6.5 with 750 mM NaCl, 75 mM sodium citrate at 42°C. Another example is use of 50% formamide, 5 x SSC (0.75 M NaCl, 0.075 M sodium citrate), 50 mM sodium phosphate (pH 6.8), 0.1% sodium pyrophosphate, 5 x Denhardt's solution, sonicated salmon sperm DNA (50 µg/ml), 0.1% SDS, and 10% dextran sulfate at 42°C, with washes at 42°C in 0.2 x SSC and 0.1% SDS.

When hK2 polypeptide is expressed in a recombinant cell other than one of human origin, the hK2 polypeptide is completely free of proteins or polypeptides of human origin. However, it is necessary to purify hK2 polypeptide from recombinant cell proteins or polypeptides to obtain preparations that are substantially homogeneous as to hK2 polypeptide. For example, the culture medium or lysate can be centrifuged to remove particulate cell debris. The membrane and soluble protein fractions are then separated. The hK2 polypeptide may then be purified from the soluble

protein fraction and, if necessary, from the membrane fraction of the culture lysate. HK2 polypeptide can then be purified from contaminant soluble proteins and polypeptides by fractionation on immunoaffinity or ion-exchange columns; ethanol precipitation; reverse phase HPLC; chromatography on silica
5 or on a cation-exchange resin such as DEAE; chromatofocusing; SDS-PAGE; ammonium sulfate precipitation; gel filtration using, for example, Sephadex G-75; or ligand affinity chromatography.

Once isolated from the resulting transgenic host cells, derivatives and variants of the hK2 polypeptide can be readily prepared. For
10 example, amides of the hK2 polypeptides of the present invention may also be prepared by techniques well known in the art for converting a carboxylic acid group or precursor, to an amide. A preferred method for amide formation at the C-terminal carboxyl group is to cleave the polypeptide from a solid support with an appropriate amine, or to cleave in the presence of an alcohol,
15 yielding an ester, followed by aminolysis with the desired amine.

Salts of carboxyl groups of the hK2 polypeptide may be prepared in the usual manner by contacting the peptide with one or more equivalents of a desired base such as, for example, a metallic hydroxide base, e.g., sodium hydroxide; a metal carbonate or bicarbonate base such as, for
20 example, sodium carbonate or sodium bicarbonate; or an amine base such as, for example, triethylamine, triethanolamine, and the like.

N-acyl derivatives of an amino group of the present polypeptides may be prepared by utilizing an N-acyl protected amino acid for the final condensation, or by acylating a protected or unprotected peptide. O-
25 acyl derivatives may be prepared, for example, by acylation of a free hydroxy peptide or peptide resin. Either acylation may be carried out using standard acylating reagents such as acyl halides, anhydrides, acyl imidazoles, and the like. Both N- and O-acylation may be carried out together, if desired. In addition, the internal hK2 amino acid sequence of Fig. 4 can be modified by
30 substituting one or two conservative amino acid substitutions for the positions specified, including substitutions which utilize the D rather than L form. The invention is also directed to variant or modified forms of the hK2 polypeptide

of Fig. 4. One or more of the residues of this polypeptide can be altered, so long as antigenic function is retained. Conservative amino acid substitutions are preferred—that is, for example, aspartic-glutamic as acidic amino acids; lysine/arginine/histidine as basic amino acids; leucine/isoleucine,

- 5 methionine/valine as hydrophobic amino acids;
serine/glycine/alanine/threonine as hydrophilic amino acids.

Acid addition salts of the polypeptides may be prepared by contacting the polypeptide with one or more equivalents of the desired inorganic or organic acid, such as, for example, hydrochloric acid. Esters of
10 carboxyl groups of the polypeptides may also be prepared by any of the usual methods known in the art.

Once isolated, hK2 polypeptide and its antigenically active variants, derivatives and fragments thereof can be used in assays for hK2 in samples derived from biological materials suspected of containing hK2 or
15 anti-hK2 antibodies, as disclosed in detail in Serial No. 08/096,946. For example, the hK2 polypeptide can be labelled with a detectable label, such as via one or more radiolabelled peptidyl residues, and can be used to compete with endogenous hK2 for binding to anti-hK2 antibodies, i.e., as a "capture antigen" to bind to anti-hK2 antibodies in a sample of a physiological fluid,
20 via various competitive immunoassay format for hK2 which uses immobilized anti-hK2 antibodies is carried out by:

- (a) providing an amount of anti-hK2 antibodies attached to a solid surface;
- (b) mixing the sample of physiological fluid to be tested with a
25 known amount of hK2 polypeptide which comprises a detectable label, to produce a mixed sample;
- (c) contacting said antibodies on said solid surface with said mixed sample for a sufficient time to allow immunological reactions to occur between said antibodies and said hK2, and between
30 said antibodies and said labelled polypeptide;
- (d) separating the solid surface from the mixed sample;

- (e) detecting or determining the presence or amount of labelled polypeptide either bound to the antibodies on the solid surface or remaining in the mixed sample; and
- (f) determining from the result in step (e) the presence or amount of said hK2 in said sample.

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In another format which can detect endogenous hK2 in a sample by a competitive inhibition immunoassay, a known amount of anti-hK2 antibody is added to a sample containing an unknown amount of endogenous hK2. The known amount is selected to be less than the amount required to complex all of the hK2 suspected to be present, e.g., that would be present in a sample of the same amount of physiological fluid obtained from a patient known to be prostate cancer. Next, a known amount of the hK2 polypeptide of the invention or a subunit thereof, comprising a detectable label is added. If endogenous hK2 is present in the sample, fewer antibodies will be available to bind the labelled hK2 polypeptide, and it will remain free in solution. If no endogenous hK2 is present, the added labelled polypeptide will complex with the added anti-hK2 antibodies to form binary complexes. Next, the binary antibody-antigen complexes are precipitated by an anti-mammal IgG antibody (sheep, goat, mouse, etc.). The amount of radioactivity or other label in the precipitate (a ternary complex) is inversely proportional to the amount of endogenous hK2 that is present in the sample, e.g., a pellet containing reduced amounts of radioactivity is indicative of the presence of endogenous hK2.

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Alternatively to the conventional techniques for preparing polyclonal antibodies or antisera in laboratory and farm animals, monoclonal antibodies against hK2 polypeptide can be prepared using known hybridoma cell culture techniques. In general, this method involves prepared an antibody-producing fused cell line, e.g., of primary spleen cells fused with a compatible continuous line of myeloma cells, and growing the fused cells either in mass culture or in an animal species from which the myeloma cell line used was derived or is compatible. Such antibodies offer many advantages in comparison to those produced by inoculation of animals, as they

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are highly specific and sensitive and relatively "pure" immunochemically. Immunologically active fragments of the present antibodies are also within the scope of the present invention, e.g., the f(ab) fragment, as are partially humanized monoclonal antibodies.

- 5 The invention will be further described by reference to the following detailed examples.

Example 1.

Construction of hK2 expression vectors

- 10 (A) Generation of recombinant baculoviruses containing pphK2 and mhK2 coding sequences

A cDNA (approximately 820 bp long) encoding the entire prepro-hK2 (pphK2) (from nucleotide #40 to #858 relative to the start site of the pphK2 transcript), as shown in Fig. 5, was synthesized from RNA of
15 human BPH tissue using reverse-transcription polymerase chain reaction (RT-PCR) technology with a pair of hK2 specific oligonucleotide primers (5'ACGCGGATCCAGCATGTGGGACCTGGTTCTCT3' SEQ ID NO: 2 and 5'ACAGCTGCAGTTTACTAGAGGTAGGGGTGGGAC 3' SEQ ID NO:3). This cDNA was generated such that 5' and 3' ends (with respect to pphK2
20 sense sequence) were bracketed with BamH1 and Pst 1 sequences respectively. The cDNA was then purified by agarose gel electrophoresis, and digested with BamH1 and Pst 1 restriction enzymes. The restricted cDNA was ligated with the BamH1-Pst 1 digested pVL1393 plasmid vector and transformed into the *E. coli* HB101 strain. *E. coli* harboring pphK2
25 cDNA/pVL1393 plasmid vector were selected and verified by restriction enzyme mapping and DNA sequencing. Plasmid pphK2 cDNA/pVL1393 was mass-produced in *E. coli* and purified by CsCl gradient ultra-centrifugation.

cDNA encoding the mature hK2 was synthesized using PCR with the aforementioned pphK2 cDNA as the template plus a pair of hK2
30 oligonucleotides (5'ACGCGGATCCAGCATGTGGGACCTGGTTCTCT3' SEQ ID NO: 2 and 5'ACCGGAATTCATGATTGTGGGAGGCTGGGAGTGT3' SEQ ID NO: 4).

As noted, the 3' end oligonucleotide was the same 3' end oligonucleotide used for synthesizing the pphK2 cDNA. However, the 5' end oligonucleotide was different from the 5' oligonucleotide used for the pphK2 cDNA, and therefore generates a cDNA coding for the mature form of hK2 (mhK2), as shown in
5 Fig. 6. The mhK2 cDNA was bracketed with EcoR1 and Pst1 sequences at the 5' and 3' ends respectively. The protein produced from the mhK2 cDNA will gain an exogenous methionine at its N-terminus. The mhK2/pVL1393 vector was generated and purified as described for pphK2/pVL1393. The DNA sequence analysis for pphK2 and mhK2 in pVL1393 showed that one
10 nucleotide (#805) has been altered (G to T) in a silent mutation.

pphK2/pVL1393 or mhK2/pVL1393 DNA (2 µg) were cotransfected with a linearized Baculogold DNA (0.5 µg; Pharmingen, San Diego, CA) into Sf9 insect cells according to Pharmingen instructions (S. Gruenwold et al., baculovirus expression vector system: Procedures and
15 Methods Manual, Pharmingen, San Diego, CA (1993)). Four to six days after the transfection, Sf9 cell spent medium containing viral particles was harvested and used to infect fresh Sf9 cells to amplify viral titers. Total RNA was isolated for Northern blot analysis of authentic pphK2 or mhK2 transcript using hK2 cDNA as a probe. Further proof of pphK2 or mhK2 transcript
20 expressed in recombinant virus infected Sf9 cells was obtained by RT-PCR and DNA sequencing. Pure recombinant baculovirus containing pphK2 or mhK2 were obtained by secondary or tertiary plaque purification protocol according to instructions from Pharmingen (S. Gruenwold et al., cited above).

25 **Example 2.**

Generation of prokaryotic expression vector

A 0.8 kb fragment representing the entire preprohK2 (pphK2) coding sequence was generated by polymerase chain reaction (PCR) using primers A (5'TATACATATGTGGGACCTGGTTCTCTCC3' SEQ ID NO.: 11)
30 and B (5'ATATGGATCCTCAGGGGTGGCTGCGATGGT3' SEQ ID NO: 12) and plasmid pVL1393 containing pphK2 as the template. The pphK2 bacterial expression vector (pBPPHK2) was prepared by standard DNA

cloning technology, (Sambrook, cited above), to subclone this 0.8 kb fragment into the NdeI/BamHI site of the plasmid pPHS579 (a gift from Dr. H. Hsiung, Eli Lilly Co, Inc.) under the control of T7 promoter. The DNA of the entire insert plus the cloning sites was sequenced to confirm that no cloning artifacts had occurred and to ensure that no anomalies in the sequence had been generated by PCR. pBPPHK2 was transformed into E. coli BL21 (DE3)Lys S (Novagen, Inc., Madison, WI).

Example 3.

10 Generation of a mammalian expression vector

To express hK2 in mammalian cell lines, a 0.8 kb fragment representing the entire preprohK2 (pphK2) coding sequence was generated by PCR using primers

A(5'ATATGGATCCATATGTCAGCATGTGGGACCTGGTTCTCTCCA3')

15 (SEQ ID NO: 17) and

B(5'ATATGGATCCTCAGGGGTTGGCTGCGATGGT3') (SEQ ID NO: 12)

and plasmid pVL1393 containing pphK2 as the template. The mammalian expression vector (pGThK2) was prepared using standard DNA cloning technology (Sambrook, 1989), to clone this 0.8 kb fragment into the BclI site of the plasmid pGT-d (a gift from Dr. Brian Grinnell, Eli Lilly, Inc.) under control of the GBMT promoter. The DNA of the entire insert plus the cloning sites was sequenced to confirm that no cloning artifacts had occurred and to ensure that no anomalies in the sequence had been generated by PCR. AV12-664 (ATCC CRL-9595), a cell line derived from a adenovirus-induced tumors in Syrian hamster, was grown in Dulbecco's modified Eagle's medium supplemented with 10% fetal bovine serum (D10F) and transfected with plasmid pGThK2 using the calcium phosphate method.

Example 4.

30 Identification of recombinant pphK2 and mhK2

A. baculovirus - insect cell system

Sf9 cells (7×10^6 /plate) were seeded onto 100 mm Corning plates with 10% fetal calf serum - Graces medium at room temperature for 1 hr. After attachment on culture plates, cells were infected with wild type or recombinant baculovirus in serum free Excell-400 medium and incubated at 27°C. Control cells were grown in the absence of virus. At designated times (24-96 hr) cells were placed in fresh Sf9-100 media deficient of either methionine or methionine and cysteine for 45-60 min at 27°C, then incubated with Promix (0.143 mCi/plate; a mixture of [35 S]-methionine and [35 S]-cysteine; 1,000-1,400 Ci/mmol; Amersham) in serum free and methionine/cysteine deficient Sf9-100 medium (Biofluids) for 5-8 hr or 20 hr. After the end of each incubation time, cells and spent media were separated by centrifugation (1,000 rpm; Beckman J-6B; Beckman, Fullerton, CA). Cells were washed and centrifuged (13,000 rpm; Biofuge 13, Baxter) twice. The washed cells were lysed by freeze/thaw in a detergent buffer (10 mM Tris, pH 7.5; 130 mM NaCl, 1% Triton X-100, 10 mM NaF; 10 mM NaPi, 10mM Nappi, pH7.5) or H₂O and centrifuged to obtain cytosol and insoluble cellular fractions. Protein contents of the above samples were determined by either the Bradford or Lowry method (BioRad, Inc., Melville, N.Y.). The above spent media, cytosol and insoluble cellular fraction were frozen and stored separately until used. A duplicate set of samples were prepared without 35 S-labeling.

For SDS-polyacrylamide gel electrophoresis (PAGE) analysis of expression of hK2 protein in Sf9 cells, samples were added to sample buffer (U.K. Laemmli, *Nature*, 227, 680 (1970)), heated at 95°C for 5 minutes and subjected to SDS-PAGE under reducing conditions.

Northern blot analysis was routinely used to screen and isolate clonal recombinant baculoviruses expressing pphK2 or mhK2 mRNA. A comparison of the corresponding lanes in both autoradiographs of the Northern blot and photographs of ethidium bromide staining of RNA shows that mRNA for pphK2 or mhK2 was present in recombinant virus infected Sf9 but not in wild type virus-infected cells. Moreover, each of the pphK2 or mhK2 mRNA positive lanes represents RNA isolated from Sf9 cells infected

with recombinant viruses derived from a single viral plaque. Thus, the results suggest that high frequency (100%) of recombinant baculovirus containing either pphK2 or mhK2 was obtained from the above cotransfection.

Furthermore, the sequences of pphK2 or mhK2 expressed in viral infected Sf9 cells were confirmed by a combination of RT-PCR, cloning and DNA sequencing.

To determine whether the pphK2 protein is expressed in the insect cell Sf9, time course studies using ³⁵S-labeling of *de novo* synthesis of protein was performed and detected by SDS denaturing polyacrylamide gel electrophoresis (PAGE). As seen in the autoradiograph (Fig. 1), a unique protein (about 28 KDa) was found in pphK2-recombinant virus-infected Sf9 cells at 35-74 hour post-infection. This band was missing in uninfected cells or cells infected with wild type virus. The viral polyhedron protein (about 32 KDa) was found (Fig. 1) as expected in Sf9 cells infected with wild type virus, whereas it was not expressed by recombinant virus (Fig. 1). The protein was detected in cytosol when subcellular fractions (cytosol vs. insoluble fraction) was prepared by lysing cells with H₂O and freeze-thaw, whereas this 28 KD protein was detected in insoluble fraction when prepared by a detergent buffer and freeze-thaw (data not shown).

The mhK2 protein was also expressed in the insect cell Sf9, ³⁵S-labeling of *de novo* synthesized protein was performed. As seen in the autoradiograph (Fig. 2), a unique protein (about 28 KDa) was found in the insoluble fraction of mhK2-recombinant virus-infected Sf9 cells at 48 hours post-infection. This band was missing in uninfected cells or cells infected with wild type virus. The viral polyhedron protein (about 32 KDa) was found in wild type virus-infected cells, whereas it was not expressed in cells infected with recombinant virus (Fig. 1). When the cytosol fraction was examined, no 28 KDa band was observed.

30 B. E. Coli system

Plasmid pBPPHK2 was transformed into *E. coli* BL21 (DE3) pLysS (Novagen, Inc., Madison, WI). This strain contains a chromosomal

copy of T7 RNA polymerase under the control of inducible LacUV5 promoter. Upon addition of IPTG (isopropyl- β -D-thiogalactopyranoside) the expression of the T7 RNA polymerase is induced which in turn activates the T7 promoter resulting in overproduction of the gene product under control of this promoter. To determine whether the product of the pphK2 gene would be expressed from pBPPHK2, single colonies of BL21 *E. Coli* transformed with pBPPHK2 were grown to O.D.₆₀₀ = 0.2 in 10 ml LB media plus ampicillin (100 μ g/ml) and induced with 0.4 mM IPTG (Sigma, Inc.). Cells were harvested 2 hours after induction by centrifugation and resuspended in 1.5 ml SDS/PAGE sample buffer (U.K. Laemmli, *Nature*, 227, 680 1970) before SDS/PAGE analysis. The cell pellet from the IPTG-induced culture was resuspended in 0.05M Tris, pH 8.0 (at 9ml/gm cell pellet) and stirred at room temperature (25°C, r.t.) for 1 hour. Lysozyme (4 mg/ml) was added to this suspension (at 1 ml/gm cell pellet) and the suspension was stirred at r.t. for 30 min followed by incubation on ice for 30 min. The suspension was sonicated for 2 min at 150 watts and centrifuged at 3000xg to isolate the inclusion bodies. Inclusion bodies were resuspended in running buffer (25 mM Tris, 192 mM glycine, 0.1% SDS) and after centrifugation both the pellet and the supernatant were analyzed by SDS/PAGE.

About 90% of the pphK2 was found to be in the supernatant fraction which indicated that pphK2 is soluble in 0.1% SDS. To prepare samples for amino acid sequence analysis, 20 μ l of inclusion body lysate was subjected to SDS/PAGE on a 4-20% gradient gel (BIO-RAD, Inc., Melville, N.Y.). The protein was blotted from the gel onto 0.2 μ PVDF paper (BIO-RAD) and stained with Coomassie blue. The protein band of interest was cut out from the blot and subjected to amino acid sequencing using a protein sequencer model 477A (Applied Biosystem, Inc., Foster City, CA).

The induced cells overproduced large amounts of a polypeptide with apparent molecular mass of about 28kd (Figure 3). Densitometric analysis indicated that this protein comprised approximately 40% of total cellular protein. The size of this protein as determined by an SDS-PAGE gel was comparable to that predicted from coding sequence for pphK2. To

confirm that this protein is pphK2, the sequence of the first 10 amino acids (MWDLVLSIAL) (SEQ ID NO: 13) from the N-terminus was determined. This sequence agrees perfectly with that deduced from the DNA sequence of pphK2 cDNA. As noted, it has different identity from the first 10 amino acids of both pphK1 (MWFLVLCLAL) (SEQ ID NO: 14) and pphK3 (MWVPVFLTL) (SEQ ID NO: 15). It also shows that this protein is not modified or processed at the N-terminus either during or after expression in *E. coli*. These results demonstrate that we were able to accurately express pphK2 in *E. coli* from pBPPHK2.

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C. Mammalian System

1. Isolation and Purification of protein

Plasmid pGThK2 was transformed into hamster cell line AV12-664 (ATCC-CRL-9595). To determine whether the product of the pphK2 gene would be expressed from pGThK2, AV12-pGThK2 #2 was grown in D10F + 200nM MTX. At about 60% confluency the cells were washed with Hank's balanced salt solution and resuspended in serum-free HH4 medium. The spent medium was collected after 7 days (serum-free spent medium) and stored at -20°C. Figure 8 depicts a SDS-PAGE confirming expression of recombinant pphK2 in a mammalian cell line. AV12-pGThK2 (Lane 4-6) and AV12-pGT-d (Lane 3) clonal cell lines were grown in D10F media. About 300µl of spent medium from the above clones were concentrated and subjected to SDS/PAGE along with See Blue MW marker (lane 1) and pphK2 lysate from *E. coli* cells (lane 2). The gel was blotted onto nitrocellulose paper and immunoblotted using a 1/1000 dilution of anti-pphK2 rabbit antiserum. HRP-goat anti-rabbit was used as the secondary probe and the blot was developed by DAB plus H₂O₂. Lane 3 (AV12-pGT-d) is AV12 transfected with vector without insert.

To purify the protein, the serum-free spent medium was concentrated from 5-10 fold by ultrafiltration with a 10 kDa molecular weight cutoff membrane then dialyzed overnight at 4°C versus 50 mM sodium bicarbonate, pH 8. Samples were filtered with 0.2 µ filters and then pumped

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directly onto a TSK DEAE-5PW HPLC column (21 mm X 150 mm) at a flow rate of 5 mL/min. Buffer A contained 50 mM sodium bicarbonate, pH 7.9. Buffer B contained 50 mM sodium bicarbonate plus 0.5 M sodium chloride, pH 7.6. The elution profile shown in Figure 9 was developed with a gradient from 0-50% Buffer B over 35 min; 50-100% B from 35-40 min and isocratic elution at 100% B for 5 min before re-equilibration in Buffer A. The flow rate was 5mL/min throughout.

DEAE fractions were assayed for the presence of hK2 by ELISA using rabbit anti-pphK2 as primary antibodies. The ELISA assayed showed a peak of hK2 activity which eluted at approximately 0-2M NaCl (shown as the triangle line in Figure 9), which correlated well with the appearance of a 34 kDa band of protein seen by SDS-PAGE in the same fractions (data not shown).

Fractions with hK2 activity were pooled and concentrated by ultrafiltration with 10 kDa membranes to approximately 5-8 mL where upon solid ammonium sulfate was added to make a final concentration of 1 M. This sample was then injected onto a PolyLC. polypropyl aspartamide column, 1000A pore size, 4.6 mm X 200 mm, to resolve protein by hydrophobic interaction chromatography (HIC, see Figure 10). Buffer A was 20 mM Na phosphate, 1.2 M Na sulfate pH 6.3 and Buffer B was 50 mM Na phosphate, 5% 2-propanol, pH 7.4. The elution gradient was 0-20% B over 5 min; 20-55% B from 5-20 min, isocratic at 55% B from 20-23 min, 55-100% B from 23-25 min; isocratic at 100% B for 2 min before re-equilibration Buffer A. The flow rate was 0.7 mL/min. Greater than 90% of the A_{280} was not retained on HIC column. The main peak retained on HIC, which eluted at 22 min, also showed the highest peak of activity by ELISA assay (triangle line, Figure 10).

HIC fractions which tested positive for hK2 on ELISA were pooled, ultrafilter concentrated as above to a volume less than 1 mL then injected on a 10/30 Pharmacia S12 size exclusion column equilibrated in 100 mM ammonium acetate. The flow rate was 0.7 mL/min. When the 22 min peak from HIC was resolved by size exclusion chromatography, typically

about 80-90% of the protein A₂₈₀ eluted at 19.4 min, a retention time consistent with a protein of approximately 34 kDa (Figure 11). The only other protein peak on SEC, eluting at 16.7 min, corresponded to an about 70KDa protein seen also in previous purification steps.

- 5 To examine the efficiency of our purification scheme, 1.5 µg of purified phK2 was subjected to SDS/PAGE in the presence or absence of β-mercaptoethanol (BME), and the gel was stained with silver. Results showed that the phK2 in our sample was about 95% pure (Fig. 12). It also showed that pro-hK2 migrated at about 30 KD in the absence of BME, and it
10 migrated at about 34 kDa in the presence of BME. This pattern is similar to that observed for the PSA purified from seminal fluid (Fig. 12).

- Recombinant phK2 is recognized by rabbit anti-pphK2, rabbit anti-PSA and a murine monoclonal antibody directed against a polypeptide covering amino acids 41-56 of hK2, when analyzed on WESTERN blots or
15 when dried down on microtiter plates. However, phK2 was not detectable by these antibodies in sandwich assays. These results further demonstrate that the phK2 and PSA are conformationally different and the antibodies currently available to PSA or hK2 can not detect phK2 in its native form. Furthermore, phK2 was not detectable by the Tandem R or free-PSA assays (immunological
20 assays for detection of PSA in serum).

A sample of the hybridoma (HK1A 523.5) secreting the murine monoclonal antibody has been deposited in the American Type Culture Collection, Rockville, MD, and assigned ATCC HB-11876.

25 2. Amino Acid Analysis and Protein Sequencing of phK2

- The peak collected off size exclusion chromatography (SEC) in ammonium acetate was lyophilized to remove the buffer then reconstituted in water. An aliquot (2.5µg) of this sample was loaded on a Poron membrane (Beckman instruments) and subjected to automated N-terminal sequence
30 analysis on an Applied Biosystems model 477A protein sequencer which yielded the following sequence:

Val-Pro-Leu-Ile-Gln-Ser-Arg-Ile-Val-Gly-Gly-Trp-Glu- (SEQ ID NO: 18). An aliquot of the same sample in water was also hydrolyzed in gaseous 6 N HCl under vacuum for 20 h at 112 °C then reconstituted in 0.1N HCl and analyzed on an Hewlett Packard Aminoquant amino acid analyzer utilizing pre-column
5 derivatization of amino acids with OPA for primary and FMOC for secondary amines.

No competing sequence was evident from the profile of amino acids released sequentially by the Edman degradation procedure. By analogy to PSA this protein is pro hK2, since the known sequence of mature PSA has
10 been shown to begin with Ileu-Val-Gly-etc and pro PSA has been postulated to have an extra 7 amino acids at the N-terminus. Amino acid analysis of this protein yielded an amino acid composition consistent with the recombinant sequence of prohK2. These results demonstrate that pphK2 was accurately expressed in the mammalian cell line AV12-664 from pGThK2.

15

Example 5.

Production of antibodies to recombinant pphK2

A. E. Coli System

To prepare pphK2 for rabbit immunization, the inclusion bodies
20 obtained from bacterial cultures of BL21 (pBpphK2) after IPTG induction as in Example 4B were resuspended in 100µl SDS/PAGE sample buffer/ml bacterial culture and electrophoresed on preparative SDS/PAGE. The pphK2 band was excised and electroeluted from the gel into running buffer (25mM Tris, 192 mM glycine, 0.1% SDS) and used as the immunogen. Two rabbits
25 were each immunized with 100µg of the immunogen in complete Freund's adjuvant and were boosted twice in three week intervals with 100µg of the immunogen in incomplete Freund's adjuvant and PBS, respectively. Rabbit anti-pphK2 sera was obtained one week following the second boost. The presence of anti-pphK2 in the rabbit antiserum was shown by ELISA (data not
30 shown). Once confirmed by this method, the highest titer antiserum was tested on Western blots using lysates from IPTG induced or non-induced cultures of BL21 (pBpphK2). It was evident that the antiserum contained

antibodies highly specific for the pphK2 protein since a protein band at about 28kd corresponding to pphK2 was recognized only in the induced lysate. The antiserum also recognized the purified pphK2 further showing the specificity of the antibodies to pphK2. The above data demonstrate that the antibodies
5 recognize the prepro-form of hK2.

To delineate if the antiserum recognizes the mature form of hK2 (mhK2), mhK2 was expressed in *E. coli* as a glutathione S-transferase fusion protein (GST-mhK2, 58kd), and the cell lysate was immunoblotted using anit-pphK2 rabbit antiserum. It was evident that anti-pphK2 antiserum
10 recognized the GST-mhK2, demonstrating that antibodies were at least in part against the mature region of pphK2. To examine the pattern recognized in seminal fluid by anti-pphK2 antibodies, seminal fluid was prepared from pooled semen as described by Sensabaugh and Blake, *J. Urology*, 149, 1523 (1990), and immunoblotted with anti-pphK2 rabbit antiserum. The antiserum
15 recognized a major band at about 34kd plus several minor bands at lower MW. The pre-immune serum did not recognize any bands in any of the above experiments, showing that the antibodies were generated by immunization.

To determine whether there are any pphK2-specific antibodies
20 in rabbit anit-pphK2 antiserum, the antibodies cross-reacting to PSA were absorbed out of the antiserum by a PSA affinity resin. Specifically, 1ml of the sera was diluted with 1mL 100 mM HEPES, pH 7.5 and incubated with native PSA-bound Affigel-10 for 3.5 hours at 4°C. The mixture was used poured into a column, the flow-through was collected and the column was
25 washed with 30 ml HEPES buffer. Antibodies bound to the column (eluate) were eluted by acetic acid (1N, pH 4.0) and neutralized to pH 6.6 with NH₄OH. Native PSA was isolated from seminal fluid as described by Sensabaugh and Blake, cited above. ppPSA was purified from *E. coli* transformed with plasmid pPHS579 (containing ppPSA under control of T7
30 promoter) using a procedure analogous to pphK2 purification.

The flow-through and the column eluate were tested for Abs recognizing pphK2, ppPSA and native PSA (PSA isolated from seminal fluid)

using Western blot analysis. It was evident that antibodies in the untreated rabbit anti-pphK2 antiserum recognized all three proteins indicating that pphK2, ppPSA and seminal fluid-PSA share some similar epitopes. However, while the column eluate contained antibodies that recognized all three protein, the flow-through contained antibodies that recognized only pphK2. This indicates that anti-pphK2 antiserum contains pphK2-specific antibodies and these antibodies can be isolated by PSA affinity absorption. This system enabled us to generate anti-pphK2 antibodies which recognize both pphK2 and mhK2. Thus, utilizing immunogenic and pure recombinant hK2 protein, generate rabbit antiserum was generated which contains pphK2-specific antibodies, providing a valuable source for generating and screening for hK2-specific monoclonal antibodies.

These examples describe the use of three heterologous expression systems (i.e. both prokaryotic and eukaryotic) for the successful expression of the hK2 polypeptide. Thus, the method of the invention enables production of large quantities of substantially pure hK2 polypeptide. The polypeptide can be used both to study its biological functions and to produce immunodetection reagents such as labelled hK2 polypeptide, labelled fragments thereof and antibodies thereto. The immunoreagents can provide a method to purify native hK2 and to study the properties of the purified native hK2 polypeptide.

The pphK2 overproduced in *E. coli* can be readily solubilized in 0.1% SDS, thus solubility is not a problem. This is in contrast to the expression of human salivary kallikrein protein, hK1, in *E. coli*, which was found in insoluble inclusion bodies (J. Wang, et al *Biochem. J.*, 276, 63 (1991)). In contrast, the present invention yields almost pure protein which can be purified to homogeneity by preparative SDS-PAGE. This purified recombinant pphK2 can be used for the generation of monoclonal and polyclonal antibodies.

As shown above, Baculogold viral DNA can be used to generate a recombinant baculovirus containing pphK2 or mhK2. Use of Baculogold viral DNA provides high selection of positive recombinant

5 baculoviruses. Indeed, Northern blot analysis showed a high frequency of recombinant virus expressing pphK2 or mhK2 mRNA. Moreover, SDS-PAGE analysis showed that both pphK2 and mhK2 recombinant viruses produced unique proteins with sizes similar to the calculated molecular weights for pphK2 or mhK2. Although the levels of the recombinant hK2 expressed in insect system may not be as high as in *E. coli*, the hK2 protein produced in baculovirus-insect system may contain the secreted form which would be more like the natural form of the protein.

10 Plasmids pphK2/pVL1393 in *E. coli* H13101 has been deposited in the American Type Culture Collection, Rockville, MD, USA on May 2, 1994 under the provisions of the Budapest Treaty and have been assigned accession number ATCC 69614.

15 The invention has been described with reference to various specific and preferred embodiments and techniques. However, it should be understood that many variations and modifications may be made while remaining within the spirit and scope of the invention.

SEQUENCE LISTING

(1) GENERAL INFORMATION:

- (i) APPLICANT: Mayo Foundation for Medical Education
and Research
Hybritech Incorporated
Tindall, Donald J.
Young, Charles Y.F.
Saedi, Mohammed S.
- (ii) TITLE OF INVENTION: Recombinant HK2 Polypeptide
- (iii) NUMBER OF SEQUENCES: 18
- (iv) CORRESPONDENCE ADDRESS:
 - (A) ADDRESSEE: Schwegman, Lundberg & Woessner, P.A.
 - (B) STREET: 3500 IDS Center
 - (C) CITY: Minneapolis
 - (D) STATE: MN
 - (E) COUNTRY: USA
 - (F) ZIP: 55402
- (v) COMPUTER READABLE FORM:
 - (A) MEDIUM TYPE: Floppy disk
 - (B) COMPUTER: IBM PC compatible
 - (C) OPERATING SYSTEM: PC-DOS/MS-DOS
 - (D) SOFTWARE: PatentIn Release #1.0, Version #1.25
- (vi) CURRENT APPLICATION DATA:
 - (A) APPLICATION NUMBER:
 - (B) FILING DATE:
 - (C) CLASSIFICATION:
- (viii) ATTORNEY/AGENT INFORMATION:
 - (A) NAME: Raasch, Kevin W.
 - (B) REGISTRATION NUMBER: 35,561
 - (C) REFERENCE/DOCKET NUMBER: 150.148WO1
- (ix) TELECOMMUNICATION INFORMATION:
 - (A) TELEPHONE: 612-339-0331
 - (B) TELEFAX: 612-339-3061

(2) INFORMATION FOR SEQ ID NO:1:

- (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 237 amino acids
 (B) TYPE: amino acid
 (C) STRANDEDNESS: single
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:

```

Ile Val Gly Gly Trp Glu Cys Glu Lys His Ser Gln Pro Trp Gln Val
1      5      10      15
Leu Val Ala Ser Arg Gly Arg Ala Val Cys Gly Gly Val Leu Val His
20     25     30
Pro Gln Trp Val Leu Thr Ala Ala His Cys Ile Arg Asn Lys Ser Val
35     40     45
Ile Leu Leu Gly Arg His Ser Leu Phe His Pro Glu Asp Thr Gly Gln
50     55     60
Val Phe Gln Val Ser Thr Ser Phe Pro His Pro Leu Tyr Asp Met Ser
65     70     75     80
Leu Leu Lys Asn Arg Phe Leu Arg Pro Gly Asp Asp Ser Ser His Asp
85     90     95
Leu Met Leu Leu Arg Leu Ser Glu Pro Ala Glu Leu Thr Asp Ala Val
100    105    110
Lys Val Met Asp Leu Pro Thr Gln Glu Pro Ala Leu Gly Thr Thr Cys
115    120    125
Tyr Ala Ser Gly Trp Gly Ser Ile Glu Pro Glu Glu Phe Leu Thr Pro
130    135    140
Lys Lys Leu Gln Cys Val Gln Leu His Val Ile Ser Asn Asp Val Cys
145    150    155    160
Ala Gln Val His Pro Gln Lys Val Thr Lys Phe Met Leu Cys Ala Gly
165    170    175
Arg Trp Thr Gly Gly Lys Ser Thr Cys Ser Gly Asp Ser Gly Gly Pro
180    185    190
Leu Val Cys Asn Gly Val Leu Gln Gly Ile Thr Ser Trp Gly Ser Glu
195    200    205
Pro Cys Ala Leu Pro Glu Arg Pro Ser Leu Tyr Thr Lys Val Val His
210    215    220
Tyr Arg Lys Trp Ile Lys Asp Thr Ile Val Ala Asn Pro
225    230    235

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32

(2) INFORMATION FOR SEQ ID NO:2:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 32 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:2:

ACGCGGATCC AGCATGTGGG ACCTGGTTCT CT

32

(2) INFORMATION FOR SEQ ID NO:3:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 33 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:3:

ACAGCTGCAG TTTACTAGAG GTAGGGGTGG GAC

33

(2) INFORMATION FOR SEQ ID NO:4:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 34 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:4:

ACCGGAATTC ATGATTGTGG GAGGCTGGGA GTGT

34

(2) INFORMATION FOR SEQ ID NO:5:

- (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 832 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(ix) FEATURE:

- (A) NAME/KEY: CDS
 (B) LOCATION: 10..792

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:5:

```

GGATCCAGC ATG TGG GAC CTG GTT CTC TCC ATC GCC TTG TCT GTG GGG      48
Met Trp Asp Leu Val Leu Ser Ile Ala Leu Ser Val Gly
   1             5             10

TGC ACT GGT GCC GTG CCC CTC ATC CAG TCT CGG ATT GTG GGA GGC TGG      96
Cys Thr Gly Ala Val Pro Leu Ile Gln Ser Arg Ile Val Gly Gly Trp
   15             20             25

GAG TGT GAG AAG CAT TCC CAA CCC TGG CAG GTG GCT GTG TAC AGT CAT     144
Glu Cys Glu Lys His Ser Gln Pro Trp Gln Val Ala Val Tyr Ser His
   30             35             40             45

GGA TGG GCA CAC TGT GGG GGT GTC CTG GTG CAC CCC CAG TGG GTG CTC     192
Gly Trp Ala His Cys Gly Gly Val Leu Val His Pro Gln Trp Val Leu
             50             55             60

ACA GCT GCC CAT TGC CTA AAG AAG AAT AGC CAG GTC TGG CTG GGT CGG     240
Thr Ala Ala His Cys Leu Lys Lys Asn Ser Gln Val Trp Leu Gly Arg
             65             70             75

CAC AAC CTG TTT GAG CCT GAA GAC ACA GGC CAG AGG GTC CCT GTC AGC     288
His Asn Leu Phe Glu Pro Glu Asp Thr Gly Gln Arg Val Pro Val Ser
             80             85             90

CAC AGC TTC CCA CAC CCG CTC TAC AAT ATG AGC CTT CTG AAG CAT CAA     336
His Ser Phe Pro His Pro Leu Tyr Asn Met Ser Leu Leu Lys His Gln
             95             100             105

AGC CTT AGA CCA GAT GAA GAC TCC AGC CAT GAC CTC ATG CTG CTC CGC     384
Ser Leu Arg Pro Asp Glu Asp Ser Ser His Asp Leu Met Leu Leu Arg
            110             115             120             125

CTG TCA GAG CCT GCC AAG ATC ACA GAT GTT GTG AAG GTC CTG GGC CTG     432
Leu Ser Glu Pro Ala Lys Ile Thr Asp Val Val Lys Val Leu Gly Leu
            130             135             140             145

CCC ACC CAG GAG CCA GCA CTG GGG ACC ACC TGC TAC GCC TCA GGC TGG     480
Pro Thr Gln Glu Pro Ala Leu Gly Thr Thr Cys Tyr Ala Ser Gly Trp
            145             150             155

GGC AGC ATC GAA CCA GAG GAG TTC TTG CGC CCC AGG AGT CTT CAG TGT     528
Gly Ser Ile Glu Pro Glu Glu Thr Leu Arg Pro Arg Ser Leu Gln Cys
            160             165             170

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34

GTG AGC CTC CAT CTC CTG TCC AAT GAC ATG TGT GCT AGA GCT TAC TCT Val Ser Leu His Leu Leu Ser Asn Asp Met Cys Ala Arg Ala Tyr Ser 175 180 185	576
GAG AAG GTG ACA GAG TTC ATG TTG TGT GCT GGG CTC TGG ACA GGT GGT Glu Lys Val Thr Glu Phe Met Leu Cys Ala Gly Leu Trp Thr Gly Gly 190 195 200 205	624
AAA GAC ACT TGT GGG GGT GAT TCT GGG GGT CCA CTT GTC TGT AAT GGT Lys Asp Thr Cys Gly Gly Asp Ser Gly Gly Pro Leu Val Cys Asn Gly 210 215 220	672
GTG CTT CAA GGT ATC ACA TCA TGG GGC CCT GAG CCA TGT GCC CTG CCT Val Leu Gln Gly Ile Thr Ser Trp Gly Pro Glu Pro Cys Ala Leu Pro 225 230 235	720
GAA AAG CCT GCT GTG TAC ACC AAG GTG GTG CAT TAC CGG AAG TGG ATC Glu Lys Pro Ala Val Tyr Thr Lys Val Val His Tyr Arg Lys Trp Ile 240 245 250	768
AAG TAC ACC ATC GCA GCC AAC CCC TGAGTGCCCC TGTCGCCACC CTACCTCTAG Lys Tyr Thr Ile Ala Ala Asn Pro 255 260	822
TAAACTGCAG	832

(2) INFORMATION FOR SEQ ID NO:6:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 261 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:6:

Met Trp Asp Leu Val Leu Ser Ile Ala Leu Ser Val Gly Cys Thr Gly 1 5 10 15
Ala Val Pro Leu Ile Gln Ser Arg Ile Val Gly Gly Trp Glu Cys Glu 20 25 30
Lys His Ser Gln Pro Trp Gln Val Ala Val Tyr Ser His Gly Trp Ala 35 40 45
His Cys Gly Gly Val Leu Val His Pro Gln Trp Val Leu Thr Ala Ala 50 55 60
His Cys Leu Lys Lys Asn Ser Gln Val Trp Leu Gly Arg His Asn Leu 65 70 75 80
Phe Glu Pro Glu Asp Thr Gly Gln Arg Val Pro Val Ser His Ser Phe 85 90 95
Pro His Pro Leu Tyr Asn Met Ser Leu Leu Lys His Gln Ser Leu Arg 100 105 110

35

Pro Asp Glu Asp Ser Ser His Asp Leu Met Leu Leu Arg Leu Ser Glu
 115 120 125

Pro Ala Lys Ile Thr Asp Val Val Lys Val Leu Gly Leu Pro Thr Gln
 130 135 140

Glu Pro Ala Leu Gly Thr Thr Cys Tyr Ala Ser Gly Trp Gly Ser Ile
 145 150 155 160

Glu Pro Glu Glu Phe Leu Arg Pro Arg Ser Leu Gln Cys Val Ser Leu
 165 170 175

His Leu Leu Ser Asn Asp Met Cys Ala Arg Ala Tyr Ser Glu Lys Val
 180 185 190

Thr Glu Phe Met Leu Cys Ala Gly Leu Trp Thr Gly Gly Lys Asp Thr
 195 200 205

Cys Gly Gly Asp Ser Gly Gly Pro Leu Val Cys Asn Gly Val Leu Gln
 210 215 220

Gly Ile Thr Ser Trp Gly Pro Glu Pro Cys Ala Leu Pro Glu Lys Pro
 225 230 235 240

Ala Val Tyr Thr Lys Val Val His Tyr Arg Lys Trp Ile Lys Tyr Thr
 245 250 255

Ile Ala Ala Asn Pro
 260

(2) INFORMATION FOR SEQ ID NO:7:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 760 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: double
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(ix) FEATURE:

- (A) NAME/KEY: CDS
- (B) LOCATION: 7..720

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:7:

GAATTC ATG ATT GTG GGA GGC TGG GAG TGT GAG AAG CAT TCC CAA CCC	48
Met Ile Val Gly Gly Trp Glu Cys Glu Lys His Ser Gln Pro	
1 5 10	
TGG CAG GTG GCT GTG TAC AGT CAT GGA TGG GCA CAC TGT GGG GGT GTC	96
Trp Gln Val Ala Val Tyr Ser His Gly Trp Ala His Cys Gly Gly Val	
15 20 25 30	
CTG GTG CAC CCC CAG TGG GTG CTC ACA GCT GCC CAT TGC CTA AAG AAG	144
Leu Val His Pro Gln Trp Val Leu Thr Ala Ala His Cys Leu Lys Lys	
35 40 45	

36

AAT AGC CAG GTC TGG CTG GGT CGG CAC AAC CTG TTT GAG CCT GAA GAC Asn Ser Gln Val Trp Leu Gly Arg His Asn Leu Phe Glu Pro Glu Asp 50 55 60	192
ACA GGC CAG AGG GTC CCT GTC AGC CAC AGC TTC CCA CAC CCG CTC TAC Thr Gly Gln Arg Val Pro Val Ser His Ser Phe Pro His Pro Leu Tyr 65 70 75	240
AAT ATG AGC CTT CTG AAG CAT CAA AGC CTT AGA CCA GAT GAA GAC TCC Asn Met Ser Leu Leu Lys His Gln Ser Leu Arg Pro Asp Glu Asp Ser 80 85 90	288
AGC CAT GAC CTC ATG CTG CTC CGC CTG TCA GAG CCT GCC AAG ATC ACA Ser His Asp Leu Met Leu Leu Arg Leu Ser Glu Pro Ala Lys Ile Thr 95 100 105 110	336
GAT GTT GTG AAG GTC CTG GGC CTG CCC ACC CAG GAG CCA GCA CTG GGG Asp Val Val Lys Val Leu Gly Leu Pro Thr Gln Glu Pro Ala Leu Gly 115 120 125	384
ACC ACC TGC TAC GCC TCA GGC TGG GGC AGC ATC GAA CCA GAG GAG TTC Thr Thr Cys Tyr Ala Ser Gly Trp Gly Ser Ile Glu Pro Glu Glu Phe 130 135 140	432
TTG GGC CCC AGG AGT CTT CAG TGT GTG AGC CTC CAT CTC CTG TCC AAT Leu Arg Pro Arg Ser Leu Gln Cys Val Ser Leu His Leu Leu Ser Asn 145 150 155	480
GAC ATG TGT GCT AGA GCT TAC TCT GAG AAG GTG ACA GAG TTC ATG TTG Asp Met Cys Ala Arg Ala Tyr Ser Glu Lys Val Thr Glu Phe Met Leu 160 165 170	528
TGT GCT GGG CTC TGG ACA GGT GGT AAA GAC ACT TGT GGG GGT GAT TCT Cys Ala Gly Leu Trp Thr Gly Gly Lys Asp Thr Cys Gly Gly Asp Ser 175 180 185 190	576
GGG GGT CCA CTT GTC TGT AAT GGT GTG CTT CAA GGT ATC ACA TCA TGG Gly Gly Pro Leu Val Cys Asn Gly Val Leu Gln Gly Ile Thr Ser Trp 195 200 205	624
GGC CCT GAG CCA TGT GCC CTG OCT GAA AAG CCT GCT GTG TAC ACC AAG Gly Pro Glu Pro Cys Ala Leu Pro Glu Lys Pro Ala Val Tyr Thr Lys 210 215 220	672
GTG GTG CAT TAC CGG AAG TGG ATC AAG TAC ACC ATC GCA GCC AAC CCC Val Val His Tyr Arg Lys Trp Ile Lys Tyr Thr Ile Ala Ala Asn Pro 225 230 235	720
TGAGTGCCCC TGTCCACCCC CTACCTCTAG TAAACTGCAG	760

37

(2) INFORMATION FOR SEQ ID NO:8:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 238 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:8:

```

Met Ile Val Gly Gly Trp Glu Cys Glu Lys His Ser Gln Pro Trp Gln
 1           5           10           15
Val Ala Val Tyr Ser His Gly Trp Ala His Cys Gly Gly Val Leu Val
          20           25           30
His Pro Gln Trp Val Leu Thr Ala Ala His Cys Leu Lys Lys Asn Ser
          35           40           45
Gln Val Trp Leu Gly Arg His Asn Leu Phe Glu Pro Glu Asp Thr Gly
          50           55           60
Gln Arg Val Pro Val Ser His Ser Phe Pro His Pro Leu Tyr Asn Met
          65           70           75           80
Ser Leu Leu Lys His Gln Ser Leu Arg Pro Asp Glu Asp Ser Ser His
          85           90           95
Asp Leu Met Leu Leu Arg Leu Ser Glu Pro Ala Lys Ile Thr Asp Val
          100          105          110
Val Lys Val Leu Gly Leu Pro Thr Gln Glu Pro Ala Leu Gly Thr Thr
          115          120          125
Cys Tyr Ala Ser Gly Trp Gly Ser Ile Glu Pro Glu Glu Phe Leu Arg
          130          135          140
Pro Arg Ser Leu Gln Cys Val Ser Leu His Leu Leu Ser Asn Asp Met
          145          150          155          160
Cys Ala Arg Ala Tyr Ser Glu Lys Val Thr Glu Phe Met Leu Cys Ala
          165          170          175
Gly Leu Trp Thr Gly Gly Lys Asp Thr Cys Gly Gly Asp Ser Gly Gly
          180          185          190
Pro Leu Val Cys Asn Gly Val Leu Gln Gly Ile Thr Ser Trp Gly Pro
          195          200          205
Glu Pro Cys Ala Leu Pro Glu Lys Pro Ala Val Tyr Thr Lys Val Val
          210          215          220
His Tyr Arg Lys Trp Ile Lys Tyr Thr Ile Ala Ala Asn Pro
          225          230          235

```

(2) INFORMATION FOR SEQ ID NO:9:

- (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 766 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

- (ix) FEATURE:
 (A) NAME/KEY: CDS
 (B) LOCATION: 1..732

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:9:

GTG	CCC	CTC	ATC	CAG	TCT	CGG	ATT	GTG	GGA	GGC	TGG	GAG	TGT	GAG	AAG	48
Val	Pro	Leu	Ile	Gln	Ser	Arg	Ile	Val	Gly	Gly	Trp	Glu	Cys	Glu	Lys	
1				5					10					15		
CAT	TCC	CAA	CCC	TGG	CAG	GTG	GCT	GTG	TAC	AGT	CAT	GGA	TGG	GCA	CAC	96
His	Ser	Gln	Pro	Trp	Gln	Val	Ala	Val	Tyr	Ser	His	Gly	Trp	Ala	His	
			20					25					30			
TGT	GGG	GGT	GTC	CTG	GTG	CAC	CCC	CAG	TGG	GTG	CTC	ACA	GCT	GCC	CAT	144
Cys	Gly	Gly	Val	Leu	Val	His	Pro	Gln	Trp	Val	Leu	Thr	Ala	Ala	His	
			35				40					45				
TGC	CTA	AAG	AAG	AAT	AGC	CAG	GTC	TGG	CTG	GGT	CGG	CAC	AAC	CTG	TTT	192
Cys	Glu	Lys	Lys	Asn	Ser	Gln	Val	Trp	Leu	Gly	Arg	His	Asn	Leu	Phe	
			50			55					60					
GAG	CCT	GAA	GAC	ACA	GGC	CAG	AGG	GTC	CCT	GTC	AGC	CAC	AGC	TTC	CCA	240
Glu	Pro	Glu	Asp	Thr	Gly	Gln	Arg	Val	Pro	Val	Ser	His	Ser	Phe	Pro	
	65				70				75						80	
CAC	CCG	CTC	TAC	AAT	ATG	AGC	CTT	CTG	AAG	CAT	CAA	AGC	CTT	AGA	CCA	288
His	Pro	Leu	Tyr	Asn	Met	Ser	Leu	Leu	Lys	His	Gln	Ser	Leu	Arg	Pro	
				85					90					95		
GAT	GAA	GAC	TCC	AGC	CAT	GAC	CTC	ATG	CTG	CTC	CGC	CTG	TCA	GAG	CCT	336
Asp	Glu	Asp	Ser	Ser	His	Asp	Leu	Met	Leu	Leu	Arg	Leu	Ser	Glu	Pro	
			100					105					110			
GCC	AAG	ATC	ACA	GAT	GTT	GTG	AAG	GTC	CTG	GGC	CTG	CCC	ACC	CAG	GAG	384
Ala	Lys	Ile	Thr	Asp	Val	Val	Lys	Val	Leu	Gly	Leu	Pro	Thr	Gln	Glu	
			115				120					125				
CCA	GCA	CTG	GGG	ACC	ACC	TGC	TAC	GCC	TCA	GGC	TGG	GGC	AGC	ATC	GAA	432
Pro	Ala	Leu	Gly	Thr	Thr	Cys	Tyr	Ala	Ser	Gly	Trp	Gly	Ser	Ile	Glu	
	130					135					140					
CCA	GAG	GAG	TTC	TTG	CGC	CCC	AGG	AGT	CTT	CAG	TGT	GTG	AGC	CTC	CAT	480
Pro	Glu	Glu	Phe	Leu	Arg	Pro	Arg	Ser	Leu	Gln	Cys	Val	Ser	Leu	His	
	145				150				155					160		
CTC	CTG	TCC	AAT	GAC	ATG	TGT	GCT	AGA	GCT	TAC	TCT	GAG	AAG	GTG	ACA	528
Leu	Leu	Ser	Asn	Asp	Met	Cys	Ala	Arg	Ala	Tyr	Ser	Glu	Lys	Val	Thr	
				165					170					175		

39

GAG TTC ATG TTG TGT GCT GGG CTC TGG ACA GGT GGT AAA GAC ACT TGT	576
Glu Phe Met Leu Cys Ala Gly Leu Trp Thr Gly Gly Lys Asp Thr Cys	
180 185 190	
GGG GGT GAT TCT GGG GGT CCA CTT GTC TGT AAT GGT GTG CTT CAA GGT	624
Gly Gly Asp Ser Gly Gly Pro Leu Val Cys Asn Gly Val Leu Gln Gly	
195 200 205	
ATC ACA TCA TGG GGC CCT GAG CCA TGT GCC CTG CCT GAA AAG CCT GCT	672
Ile Thr Ser Trp Gly Pro Glu Pro Cys Ala Leu Pro Glu Lys Pro Ala	
210 215 220	
GTG TAC ACC AAG GTG GTG CAT TAC CGG AAG TGG ATC AAG TAC ACC ATC	720
Val Tyr Thr Lys Val Val His Tyr Arg Lys Trp Ile Lys Tyr Thr Ile	
225 230 235 240	
GCA GCC AAC CCC TGAGTGCCCC TGTCCCACCC CTACCTCTAG TAAA	766
Ala Ala Asn Pro	

(2) INFORMATION FOR SEQ ID NO:10:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 244 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:10:

Val	Pro	Leu	Ile	Gln	Ser	Arg	Ile	Val	Gly	Gly	Trp	Glu	Cys	Glu	Lys
1				5					10					15	
His	Ser	Gln	Pro	Trp	Gln	Val	Ala	Val	Tyr	Ser	His	Gly	Trp	Ala	His
		20						25					30		
Cys	Gly	Gly	Val	Leu	Val	His	Pro	Gln	Trp	Val	Leu	Thr	Ala	Ala	His
		35					40					45			
Cys	Leu	Lys	Lys	Asn	Ser	Gln	Val	Trp	Leu	Gly	Arg	His	Asn	Leu	Phe
	50					55					60				
Glu	Pro	Glu	Asp	Thr	Gly	Gln	Arg	Val	Pro	Val	Ser	His	Ser	Phe	Pro
	65				70					75					80
His	Pro	Leu	Tyr	Asn	Met	Ser	Leu	Leu	Lys	His	Gln	Ser	Leu	Arg	Pro
				85					90					95	
Asp	Glu	Asp	Ser	Ser	His	Asp	Leu	Met	Leu	Leu	Arg	Leu	Ser	Glu	Pro
		100						105					110		
Ala	Lys	Ile	Thr	Asp	Val	Val	Lys	Val	Leu	Gly	Leu	Pro	Thr	Gln	Glu
		115					120					125			
Pro	Ala	Leu	Gly	Thr	Thr	Cys	Tyr	Ala	Ser	Gly	Trp	Gly	Ser	Ile	Glu
	130					135					140				
Pro	Glu	Glu	Phe	Leu	Arg	Pro	Arg	Ser	Leu	Gln	Cys	Val	Ser	Leu	His
145					150					155					160

40

Leu Leu Ser Asn Asp Met Cys Ala Arg Ala Tyr Ser Glu Lys Val Thr
 165 170 175
 Glu Phe Met Leu Cys Ala Gly Leu Trp Thr Gly Gly Lys Asp Thr Cys
 180 185 190
 Gly Gly Asp Ser Gly Gly Pro Leu Val Cys Asn Gly Val Leu Gln Gly
 195 200 205
 Ile Thr Ser Trp Gly Pro Glu Pro Cys Ala Leu Pro Glu Lys Pro Ala
 210 215 220
 Val Tyr Thr Lys Val Val His Tyr Arg Lys Trp Ile Lys Tyr Thr Ile
 225 230 235 240
 Ala Ala Asn Pro

(2) INFORMATION FOR SEQ ID NO:11:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 28 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:11:

TATACATATG TGGGACCTGG TTCTCTCC

28

(2) INFORMATION FOR SEQ ID NO:12:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 31 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:12:

ATATGGATCC TCAGGGGTTG GCTGCGATGG T

31

(2) INFORMATION FOR SEQ ID NO:13:

- (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 10 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:13:

Met Trp Asp Leu Val Leu Ser Ile Ala Leu
1 5 10

(2) INFORMATION FOR SEQ ID NO:14:

- (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 10 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:14:

Met Trp Phe Leu Val Leu Cys Leu Ala Leu
1 5 10

(2) INFORMATION FOR SEQ ID NO:15:

- (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 10 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:15:

Met Trp Val Pro Val Val Phe Leu Thr Leu
1 5 10

(2) INFORMATION FOR SEQ ID NO:16:

- (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 237 amino acids
 (B) TYPE: amino acid
 (C) STRANDEDNESS: single
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:16:

```

Ile Val Gly Gly Trp Glu Cys Glu Lys His Ser Gln Pro Trp Gln Val
1           5           10           15
Ala Val Trp Ser His Gly Trp Ala His Cys Gly Gly Val Leu Val His
20           25           30
Pro Gln Trp Val Leu Thr Ala Ala His Cys Leu Lys Lys Asn Ser Gln
35           40           45
Val Trp Leu Gly Arg His Asn Leu Phe Glu Pro Glu Asp Thr Gly Gln
50           55           60
Arg Val Pro Val Ser His Ser Phe Pro His Pro Leu Tyr Asn Met Ser
65           70           75           80
Leu Leu Lys His Gln Ser Leu Arg Pro Asp Glu Asp Ser Ser His Asp
85           90           95
Leu Met Leu Leu Arg Leu Ser Glu Pro Ala Lys Ile Thr Asp Val Val
100          105          110
Lys Val Leu Gly Leu Pro Thr Gln Glu Pro Ala Leu Gly Thr Thr Cys
115          120          125
Tyr Ala Ser Gly Trp Gly Ser Ile Glu Pro Glu Glu Phe Leu Arg Pro
130          135          140
Arg Ser Leu Gln Cys Val Ser Leu His Leu Leu Ser Asn Asp Met Cys
145          150          155          160
Ala Arg Ala Tyr Ser Glu Lys Val Thr Glu Phe Met Leu Cys Ala Gly
165          170          175
Leu Trp Thr Gly Gly Lys Asp Thr Cys Gly Gly Asp Ser Gly Gly Pro
180          185          190
Leu Val Cys Asn Gly Val Leu Gln Gly Ile Thr Ser Trp Gly Pro Glu
195          200          205
Pro Cys Ala Leu Pro Glu Lys Pro Ala Val Tyr Thr Lys Val Val His
210          215          220
Tyr Arg Lys Trp Ile Lys Asp Thr Ile Ala Ala Asn Pro
225          230          235

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43

(2) INFORMATION FOR SEQ ID NO:17:

- (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 42 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:17:

ATATGGATCC ATATGTCAGC ATGTGGGACC TGGTCTCTC CA

42

(2) INFORMATION FOR SEQ ID NO:18:

- (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 13 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:18:

Val Pro Leu Ile Gln Ser Arg Ile Val Gly Gly Trp Glu
1 5 10

WHAT IS CLAIMED IS:

1. Isolated, substantially homogeneous pre-pro hK2 polypeptide.
- 5 2. Isolated, substantially homogenous pro hK2 polypeptide.
3. Isolated, substantially homogenous mature hK2 polypeptide.
4. An antibody that is capable of specifically binding the hK2 polypeptide
10 of claims 1, 2 or 3 and which does not bind to hK3.
5. The antibody of claim 4 which is a monoclonal antibody.
6. A hybridoma cell line producing the antibody of claim 5.
- 15 7. An isolated nucleic acid molecule encoding the polypeptides of claim 1, 2 or 3.
8. An isolated nucleic acid molecule selected from the group consisting
20 of
 - (a) cDNA comprising the nucleotide sequence of the coding region of the hK2 gene;
 - (b) DNA capable of hybridizing under stringent conditions to a nucleotide sequence complementary to the nucleotide sequence
25 of (a); and
 - (c) a genetic variant of any of the DNA of (a) and (b) which encodes a polypeptide possessing an antigenic function of naturally occurring hK2 polypeptide.
- 30 9. The nucleic acid molecule of claim 8 further comprising a promoter operably linked to the nucleic acid molecule.

10. A chimeric expression vector comprising the nucleic acid molecule of claim 7 operably linked to control sequences recognized by a host cell transformed with the vector.
- 5 11. The vector of claim 10 wherein the host cell is E. coli.
12. The vector of claim 10 wherein the host cell is a mammalian cell.
13. A host cell transformed with the vector of claim 10.
- 10 14. The host cell of claim 13 which is E. coli.
15. The host cell of claim 13 which is mammalian.
- 15 16. A method of using a nucleic acid molecule encoding a hK2 polypeptide comprising expressing the nucleic acid molecule of claim 7 in a cultured host cell stably transformed with a chimeric vector comprising said nucleic acid molecule operably linked to control sequences recognized by the host cell transformed with the vector, and
20 recovering hK2 polypeptide from the host cell.
17. The method of claim 16 wherein the host cell is E. coli.
18. The method of claim 16 wherein the host cell is mammalian.
- 25 19. The method of claim 16 wherein the nucleic acid molecule is DNA.
20. The method of claim 16 wherein the hK2 polypeptide is recovered from the host cell culture medium.

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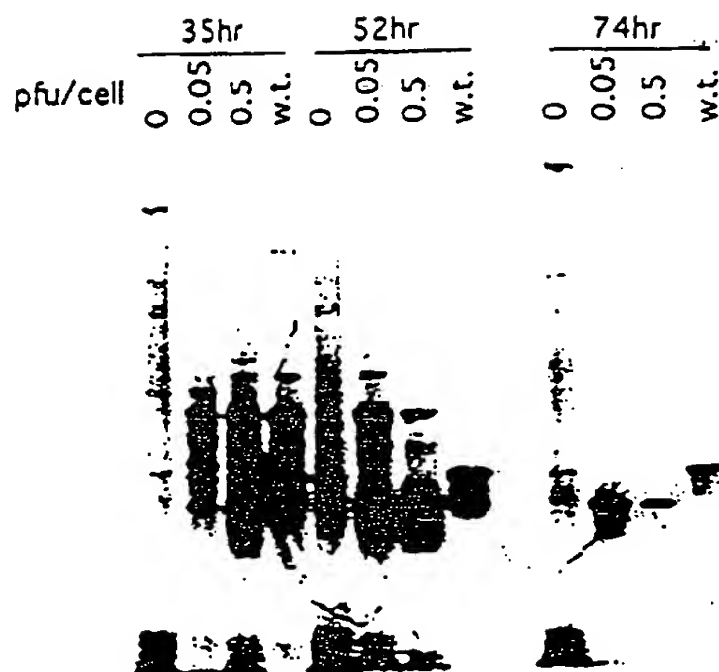


Figure 1

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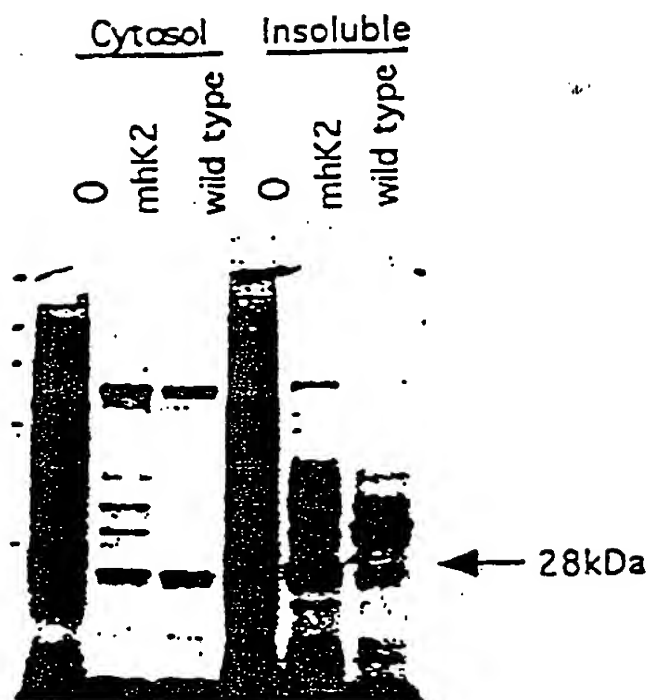
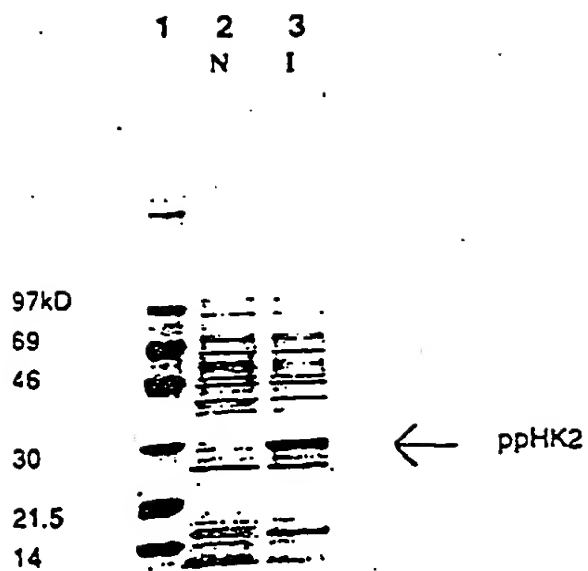


Figure 2

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Lane 1 Molecular Weight marker
2 Not - induced (N)
3 Induced (I)

Figure 3

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HK2: IVGGWECEKHSQPWQVAVWSHGWAHCGGVLVHPQWVLTA⁴¹AHCLKKNSQVWLGRHN
 HK3: "L"A"R"R"V"IRNK"VIL""S

⁵⁶
 LFEPEDTGQRVFVSHSFPHPLYNMSLLKHQSLRPDEDSSHDLMLRLSEPAKIT
 "H""VFQ""T""D""NR""GD""EL"

110 ¹⁵³
 DVVKVLGLPTQEPALGTTTCYASGWGSIEPEEFLRPRSLQCVSLHLLSNDMCA
 "A""MD""TPKK""Q""VI""V"

162 167
 RAYSEKVTEFMLCAGLWTGGKDTCCGDSGGPLVCNGVLQGITSWGPEPCALPEKP
 QVHPQ""K""R""S""S""S""R"

217 237
 AVYTKVVHYRKWIKDTIAANP
 SL""V""

Figure 4

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BamHI ← → -pphK2 → phK2
 1 GGATCCAGCATGTGGGACCTGTTCTCTCCATCGCCTTGTCTGTGGGCTGCACTTGTGCCGAGCC
 CCTAGGTCGTACACCTGGACCAAGAGAGGTAGCGGACAGACACCCACGTGACCACGGCACGG
 1▶ Met Trp Asp Leu Val Leu Ser Ile Ala Leu Ser Val Gly Cys Thr Gly Ala Val Pr
 66 CCTCATCCAGTCTCCGATTGTGGAGGCTGGGAGTGTGAGAAGCAATCCCAACCCCTGGCAGGTGG
 GGAGTAGGTCAGAGCCTAACACCCCTCCGACCCACACTCTTCGTAGGGTTGGACCGTCCACC
 19▶ oLeu Ile Gl n Ser Arg Ile Val Gly Gly Trp Gl u Cys Gl u Lys His Ser Gl n Pro Trp Gl n Val A
 131 CTGGGTACAGTCATGGATGGGCACACTGAGGGGTGTCTCTGCTGCACCCCACTGGGTGCTCACA
 GACACATGTCACTAGCTACCCGTGTGACACCCCAAGGACACCGTGGGGGTCACTCCACAGTGT
 41▶ Ia Val Tyr Ser His Gly Trp Ala His Cys Gly Gly Val Leu Val His Pro Gl n Trp Val Leu Thr
 196 GCTGCCCATTCGCTAAAGAAGATAGCCAGGTCTGGCTGGGTCCGGCACAACCTGTTTGAGCCTGA
 CGACGGGTAAACGATTCTCTTATCGGTCCAGACCGACCCAGCCGTGTTGGACAACTCCGACT
 63▶ Ala Ala His Cys Leu Lys Lys Asn Ser Gl n Val Trp Leu Gly Arg His Asn Leu Phe Gl u Pro Gl
 261 AGACACAGGGCCAGGGTCCCTGTCAAGCCACAGCTTCCACACCCCGCTCTACAATATGAGCCTTC
 TCTGTCTCCGGTCTCCAGGACAGTGGGTCTCGAAGGGTGTGGGCGAGATGTTATACTCGGAAG
 84▶ u Asp Thr Gly Gl n Arg Val Pro Val Ser His Ser Phe Pro His Pro Leu Tyr Asn Met Ser Leu L
 326 TGAAGCATCAAAAGCCTTAGACACAGATGAAGACTCCACCCATGACCTCATGCTGCTCCGCTGTCA
 ACTTCGTAGTTTCCGAATCTGCTACTCTCTGAGGTCTGCTACTGGAGTACGACGAGGCGGACAGT
 106▶ eu Lys His Gl n Ser Leu Arg Pro Asp Gl u Asp Ser Ser His Asp Leu Met Leu Leu Arg Leu Ser
 391 GAGCCTGCCAAGATCAGAGATTTGTGAAGGTCTGGGCTGCCCCACCCAGAGGCCAGCACTGGG
 CTCGGACGGTTCTAGTGTCTACAACACTCCACAGCCGGACGGGTGGGTCTCTGGTCTGTGACCC
 128▶ Gl u Pro Ala Lys Ile Thr Asp Val Val Lys Val Leu Gly Leu Pro Thr Gl n Gl u Pro Ala Leu Gl
 456 GACCACCTGCTACCTCTAGGCTGGGGCAGCATCGAACCCAGAGGATTTCTTGGCCCCAGGAGTC
 CTGGTGGACGATGCGAGTCCGACCTCTCTGCTAGCTTGTCTCTCTCAAGAACCGGGGTCTCTCAG
 149▶ y Thr Thr Cys Tyr Ala Ser Gly Trp Gly Ser Ile Gl u Pro Gl u Gl u Phe Leu Arg Pro Arg Ser L
 521 TTCAGTGTGTGAGCCTCCATCTCTGTCCAAATGACATGTGTGCTAGAGCTTACTCTGAGAAGGTG
 AAGTCACACACTCGGAGGTAGAGACAGGTTACTGTACACACGATCTCGAATGAGACTCTTCCAC
 171▶ eu Gl n Cys Val Ser Leu His Leu Leu Ser Asn Asp Met Cys Ala Arg Ala Tyr Ser Gl u Lys Val
 586 ACACAGTTCATGTTGTGTGCTGGGCTCTGGACAGTGGTAAAGACACTTGTGGGGGTGATTCTGG
 TGTCTCAAGTACACACAGACCCGAGACCTGTCCACCATTTCTGTGAACACCCCCACTAAGACC
 193▶ Thr Gl u Phe Met Leu Cys Ala Gly Leu Trp Thr Gly Gly Lys Asp Thr Cys Gly Gly Asp Ser Gl
 651 GGGTCCACTTGTCTGTAAATGGTGTGTTCAAGGTATCACAATCATGGGGCCCTGAGCCATGTGCCC
 CCCAGGTGAACAGACATTACCACCGAAGTTCCATAGTGTAGTACCCCGGACTCGGTACACGGG
 214▶ y Gly Pro Leu Val Cys Asn Gly Val Leu Gl n Gly Ile Thr Ser Trp Gly Pro Gl u Pro Cys Ala L
 716 TGCCTGAAGAGCCTGCTGTGTACACCAAGGTGGTGCATTACCGAAGTGGATCAAGTACACCATC
 ACGGACTTTTCGAGACACATGTGGTTCCACCAAGTAAATGGCTTCACTAGTTCAATGTGGTAG
 236▶ eu Pro Gl u Lys Pro Ala Val Tyr Thr Lys Val Val His Tyr Arg Lys Trp Ile Lys Tyr Thr Ile
 781 GCAGGCCAACCCCTGAGTGGCCCTGTCCACCCCTACCTCTAGTAACTGCAG
 CTTCCCTTGGGACTCAGCGGACAGGGTGGGATGAGATCATTTGACGTC
 258▶ Ala Ala Asn Pro → PstI

Figure 5

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1 GAATTCATGATTGTGGAGGCTGGGAGTGTGAGAGCATTCCCAACCC
 CTTAAATCTAACACCCCTCCGACCCCTCAGACCTTTTCGTAGGXTTGGG
 1▶ Met Ile Val Gly Gly Trp Glu Cys Glu Lys His Ser Glu Pro
 49 TGGCAGGTGKTNVTGTACAGTCAATGATAGGCACACTNTGGGGTNTTC
 ACCGTCCACCGACACATGTCACTACCTACCCGTGTGACACCCCCACAG
 15▶ Trp Glu Val Ala Val Tyr Ser His Gly Trp Ala His Cys Gly Gly Val
 97 CTGGTGCACCCCACTGGGTGCTCAGAGCTGCCCATTTGCCTAAGAGAG
 GACCAAGTGGGGTCAACCCACAGTGTGAGAGGTAACGGATTTCTTC
 31▶ Leu Val His Pro Glu Trp Val Leu Thr Ala Ala His Cys Leu Lys Lys
 145 AATAGCCAGGCTCTGGCTGGGTGGGACAACTTCTTGAGCCTGAGAGC
 TTATCGGTCCAGACCGACCCAGCCGTGTGTGGACAACTCGGACITCTG
 47▶ Asn Ser Glu Val Trp Leu Gly Arg His Asn Leu Phe Glu Pro Glu Asp
 193 ACAGCCACAGAGGGTCCCTGTCAAGCCACAGCTTCCACACCCCGCTCTAC
 TGTCCGCTCTCCACGGGACAGTCCGTGTGAGAGGGTGTGGCCAGCTG
 63▶ Thr Gly Glu Arg Val Pro Val Ser His Ser Phe Pro His Pro Leu Tyr
 241 AATATGAGCCTTCTGAAGCAGCAAAAGCCCTAGACCAAGTGAAGACCTCC
 TTATCTCGGAGAGCTTCTAGTTCGGAACTCTGGTCTACTCTCTGAGG
 79▶ Asn Met Ser Leu Leu Lys His Glu Ser Leu Arg Pro Asp Glu Asp Ser
 289 AGCCATGACCTCATGCTGCTTCCGCTTTCAGAGCCTTCCCAAGATCACA
 TCGGTACTGCACTACGACGAGGCGGACAGTCTCGGACCGGTCTAGTGT
 95▶ Ser His Asp Leu Met Leu Leu Arg Leu Ser Glu Pro Ala Lys Ile Thr
 337 GATGTTGTGAAGGTCTCTGGGCTGCCACCCAGGAGCCAGCACTGGGG
 CTACACACTTCCAGGACCCGAGCGGTGGGTCTCTGGTCTGTGACCC
 111▶ Asp Val Val Lys Val Leu Gly Leu Pro Thr Glu Glu Pro Ala Leu Gly
 385 ACCACTCTCTACGCTCAGGCTGGGCGCAGCATCGAACCAAGAGGAGTTC
 TGGTGGACGATCCGAGTCCGACCCCGTGGTAGCTTGGTCTCTCTCAG
 127▶ Thr Thr Cys Tyr Ala Ser Gly Trp Gly Ser Ile Glu Pro Glu Glu Phe
 433 TTGCGGCCCCAGGAGTCTTCACTGTGTGAGCCTCCATCTCTGTCCAA
 AACCGGGGGTCTCAGAAGTCACACACTCGGAGGTAGAGGACAGGTTA
 143▶ Leu Arg Pro Arg Ser Leu Glu Cys Val Ser Leu His Leu Leu Ser Asn
 481 GACATGTGTGCTAGAGCTTACTCTGAGAGGGTGACAGGTTTCATGTTG
 CTGTACACACGATCTCGAATGAGACTCTTCCACTGTCTCAAGTACAA
 159▶ Asp Met Cys Ala Arg Ala Tyr Ser Glu Lys Val Thr Glu Phe Met Leu
 529 TGTGCTGGGGCTCTGGACAGGTGGTAAAGACACTGTGGGGGTGATCT
 ACACGACCCGAGACCTGTCCACCATTTCTGTGACACCCCCACTAAGA
 175▶ Cys Ala Gly Leu Trp Thr Gly Gly Lys Asp Thr Cys Gly Gly Asp Ser
 577 GGGGGTCCACTTGTCTGTAATGGTGTGCTTCAAGGTATCAGATCATGG
 CCCCCAGGTGAACAGACATTACCAACAGGATTTCATAGTGTAGTACC
 191▶ Gly Gly Pro Leu Val Cys Asn Gly Val Leu Glu Gly Ile Thr Ser Trp
 625 GGCCCTGAGCCATGTGCCCTGCTGAAAAGCCTGCTGTGTACACCAAG
 CCGGACTCGGTACACGGGACCGACTTTTCCGACGACACATGTGGTTC
 207▶ Gly Pro Glu Pro Cys Ala Leu Pro Glu Lys Pro Ala Val Tyr Thr Lys
 673 GTGGTGCAATTACCGGAAGTGGATCAAGTACACCATCGGACCCCAACCC
 CACACAGTAATGGCCTTCACTAGTTCATGTGGTAGCGTCCGGTTGGGG
 223▶ Val Val His Tyr Arg Lys Trp Ile Lys Tyr Thr Ile Ala Ala Asn Pro
 721 TTAGTGCCCCCTGTCCACCCCTACCTCTAGTAACCTCCAG
 ACTCACGGGGACAGGGTGGGTATGGAGATCATTTCACGTC

Figure 6

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1 GTGCCCTCATCCAGTCTCGGATTGTGGGAGGCTGGGAGTGTGAGAAGCAATCCCAACCC
 CACGGGGAGTAGGTCAAGCCTAACACCTCCGACCCTCACACTCTTCGTAAAGGGTTGGG
 1▶ Val ProLeuIleGlnSerArgIleVal GlyGlyTrpGluCysGluLysHisSer GlnPro
 61 TGGCAGGTGGCTGTGTACAGTCATGGATGGGCACACTGTGGGGGTGTCTGGTGCACCCC
 ACCGTCCACCGACACATGTCACTACCTACCCGTGTGACACCCCCACAGGACCAAGTGGG
 21▶ TrpGlnValAlaValTyrSerHisGlyTrpAlaHisCysGlyGlyValLeuValHisPro
 121 CAGTGGGTGTCTACAGCTGCCATTGCCCTAAAGAAGATAGCCAGGTCTGGCTGGGTGGG
 GTCACCCACGAGTGTCCGACGGTAACGGATTTCCTCTATCGGTCCAGACCGACCCAGCC
 41▶ GlnTrpValLeuThrAlaAlaHisCysLeuLysLysAsnSerGlnValTrpLeuGlyArg
 181 CACAACCTGTTTGAGCCTGAAGACACAGGCCAGAGGGTCCCTGTGAGCCACAGCTTCCCA
 GTGTTGGACAAACTCGACTTCTGTGTCCGGTCTCCACGGACAGTCCGGATCCGAAGGGT
 61▶ HisAsnLeuPheGluProGluAspThr GlyGlnArgValProValSerHisSerPhePro
 241 CACCCGCTCTACAATATGAGCCTTCTGAAGCATCAAGCCTTAGACCAGATGAAGACTCC
 GTGGGCGAGATGTTATACTCGGAAGACTTCGTAGTTTCGGAATCTGGTCTACTTCTGAGG
 81▶ HisProLeuTyrAsnMetSerLeuLeuLysHisGlnSerLeuArgProAspGluAspSer
 301 AGCCATGACCTCATGTCTCCGCTGTGAGCCCTGSCAAGATCACAGTGTGTGTGAG
 TCGGTACTGGAGTACGACGAGGCGGACAGTCTCGGACGGTCTAGTGTCTACACACTTC
 101▶ SerHisAspLeuMetLeuLeuArgLeuSerGluProAlaLysIleThrAspValValLys
 361 GTCTGTGGCCTGCCACCCAGGAGCCAGCACTGGSGACCACCTGCTACGCCCTCAGGCTGG
 CAGGACCCGGACGGGTGGTCTCTCGGTCTGTACCCCTGGTGGACGATGCGGAGTCCGACC
 121▶ ValLeuGlyLeuProThrGlnGluProAlaLeuGlyThrThrCysTyrAlaSerGlyTrp
 421 GCGAGCATCGAACCCAGAGGAGTCTCTCCGCCCCAGGAGTCTTCAGTGTGTGAGCCTCCAT
 CCGTCTGATGTTGGTCTCTCTCAAGAACCGGGGTCTCTAGAGTCTACACCTCTGGAGTA
 141▶ GlySerIleGluProGluGluPheLeuArgProArgSerLeuGlnCysValSerLeuHis
 481 CTCCTGTCTCAATGACATGTGTGTAGAGCTTACTCTGAGAAGGTGACAGAGTTCATGTTG
 GAGGACAGGTTACTGTACACAGATCTCGAATGAGACTCTTCCACTGTCTCAAGTACAAC
 161▶ LeuLeuSerAsnAspMetCysAlaArgAlaTyrSerGluLysValThrGluPheMetLeu
 541 TGTGCTGGGCTCTGGACAGGTGTAAAGACACTTGTGGGGGTGATTCTGGGGGTCCACTT
 ACACGACCCGAGACCTGTCCACCATTTCTGTGAACACCCCCACTAAGACCCCCAGGTGAA
 181▶ CysAlaGlyLeuTrpThrGlyGlyLysAspThrCysGlyGlyAspSerGlyGlyProLeu
 601 GTCTGTAAATGGTGTGCTTCAAGGTATCACATCATGGGGCCCTGAGCCATGTGCCCTGCCCT
 CAGACATTACCAACGAAGTTCCATAGTGTAGTACCCCGGACTCGGTACACGGGACCGGA
 201▶ ValCysAsnGlyValLeuGlnGlyIleThrSerTrpGlyProGluProCysAlaLeuPro
 661 GAAAGCCTGCTGTGTACACCAAGGTGGTGCATTACCGGAAGTGGATCAAGTACACCATC
 CTTTTCGGACGACACATGTGGTTCCACCACGTAAATGGCCTTCACCTAGTTTCATGTGGTAG
 221▶ GluLysProAlaValTyrThrLysValValHisTyrArgLysTrpIleLysTyrThrIle
 721 GCAGCCAACCCCTGAGTGCCTCTGTCCACCCCTACCTCTAGTAAA
 CGTCGGTTGGGACTCACGGGACAGGGTGGGATGGAGATCATT
 241▶ AlaAlaAsnPro

Figure 7

hK2 Expression in AV12 Cells

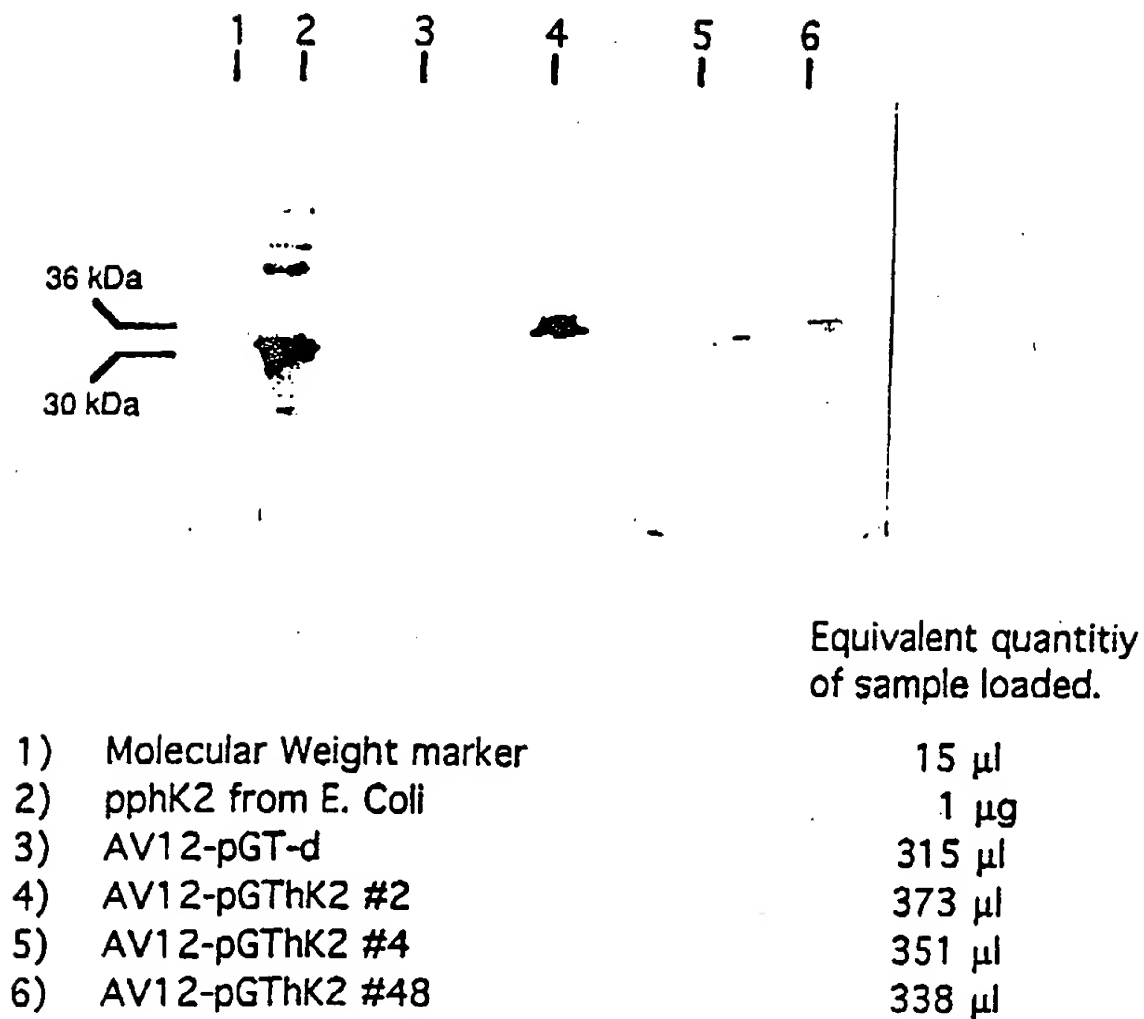


Figure 8

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DEAE Chromatography

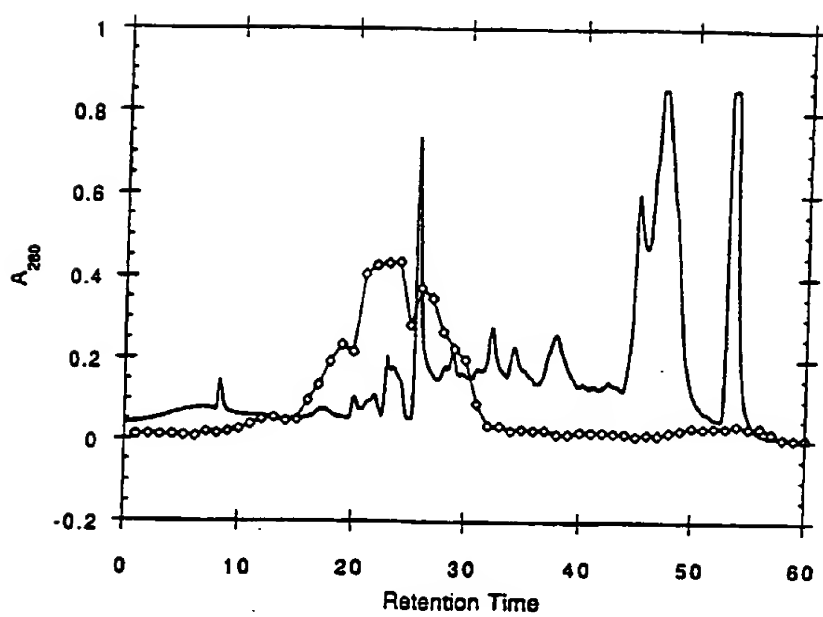


Figure 9

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Hydrophobic Interaction Chromatography

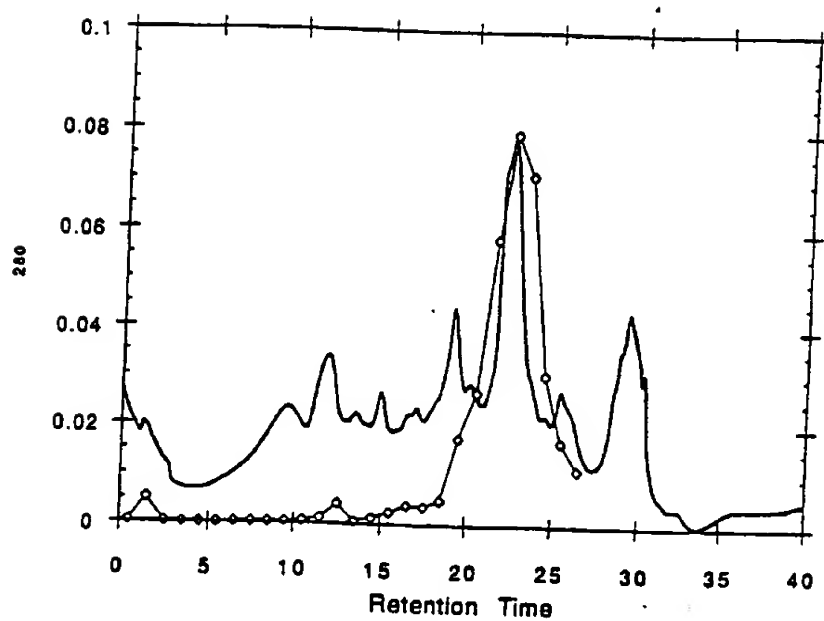


Figure 10

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Size Exclusion Chromatography

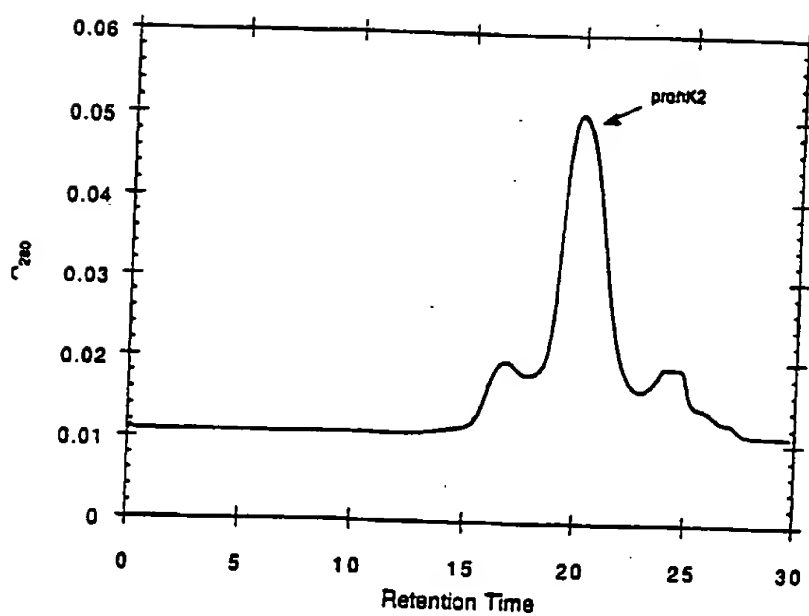
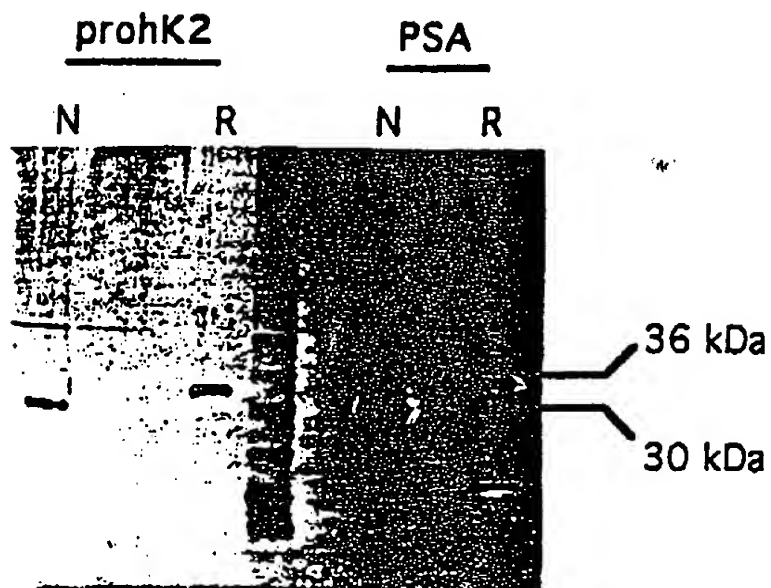


Figure 11

SDS / PAGE Analysis of prohK2 and PSA



— 1.5 μ g protein per well

Figure 12

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 95/06157

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C12N15/57 C12N9/64 C07K16/40 C12N5/20 C12N5/10
C12N1/21

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C12N C07K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	JOURNAL OF HYPERTENSION, vol. 6, no. s4, 1988 pages s395-s398, L. SCHEDLICH ET AL 'Kallikrein genes: cloning in man and expression in rat renal hypertension' see the whole document ---	1-20
X	MOLECULAR AND CELLULAR ENDOCRINOLOGY, vol. 76, 1991 pages 181-190, P. RIEGMAN ET AL 'Identification and androgen-regulated expression of two major human glandular kallikrein-1 (hGK++) mRNA species' see the whole document ---	1-20
-/--		

☒ Further documents are listed in the continuation of box C.

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"&" document member of the same patent family

Date of the actual completion of the international search

25 September 1995

Date of mailing of the international search report

13. 10. 95

Name and mailing address of the ISA

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Fax (+ 31-70) 340-3016

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Van der Schaal, C

INTERNATIONAL SEARCH REPORT

International Application No.

PCT/US 95/06157

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	DNA, vol. 6, no. 5, 1987 pages 429-437, L. SCHEDLICH ET AL 'Primary structure of a human glandular kallikrein gene' see the whole document ---	1-20
X	FEBS, vol. 236, no. 1, August 1988 pages 205-208, P. CHAPDELAINE ET AL 'High level expression in the prostate of a human glandular kallikrein mRNA related to prostate-specific antigen' see the whole document ---	1-20
A	EP-A-0 297 913 (AMGEN INC) 4 January 1989 see the whole document ---	8-20
P,X	WO-A-95 03334 (MAYO FOUNDATION) 2 February 1995 see the whole document ---	1-20
P,X	MOL. CELL. ENDOCRINOL. (1995), 109(2), 237-41 CODEN: MCEND6; ISSN: 0303-7207, 1995 SAEDI, MOHAMMAD S. ET AL 'Overexpression of a human prostate-specific glandular kallikrein, hK2, in E. coli and generation of antibodies' see the whole document -----	1-20

INTERNATIONAL SEARCH REPORT

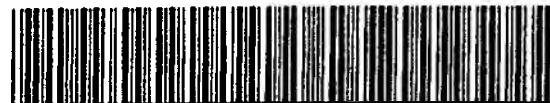
International Application No.

PCT/US 95/06157

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP-A-0297913	04-01-89	AT-T- 118547	15-03-95
		AU-B- 2075588	30-01-89
		DE-D- 3853023	23-03-95
		DE-T- 3853023	08-06-95
		ES-T- 2068201	16-04-95
		JP-T- 1503679	14-12-89
		PT-B- 87887	31-05-94
		WO-A- 8900192	12-01-89
WO-A-9503334	02-02-95	AU-B- 7252594	20-02-95

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VERSION*CORRECTED
VERSION**

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International Bureau

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(51) International Patent Classification 6: C12N 15/57, 9/64, C07K 16/40, C12N 5/20, 5/10, 1/21	A1	(11) International Publication Number: WO 95/30758 (43) International Publication Date: 16 November 1995 (16.11.95)
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(74) Agent: RAASCH, Kevin, W.; Schwegman, Lundberg & Woessner, 3500 IDS Center, 80 South Eighth Street, Minneapolis, MN 55402 (US).

(81) Designated States: AU, CA, HU, JP, KR, MX, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).

Published*With international search report.**Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.**With an indication in relation to a deposited microorganism furnished under Rule 13bis separately from the description.**Date of receipt by the International Bureau:**29 June 1995 (29.06.95)*(54) Title: **RECOMBINANT HK2 POLYPEPTIDE**

41
hK2: **IVGGWECEKHSQPWQVAVVWSHGWAHCGGVLVHPQWVLTAAHCLKNSQVWLGRHN**
hK3: *******LAAARRRV*****IRNKVIL*****S**

56
LFEPEDTGQRPVPSHSFPHPLYNMSLLKHQSLRPDEDSSHDLMLRLSEPAKIT
*******VFQ*****T*****NR*****GD*****EL*******

110 153
DVVKVLGLPTQEPALGTTTCYASGWSIEPEEFLRPSLQCVSLHLLSNDHCA
*******ND*****TPKK*****Q*****VI*****V*******

162 167
RAYSEKVTETMLCAGLWTGGKDTCCGDSGGPLVCNGVLQGITSWGPEPCALPEKP
QVHRQ***K*****R*****S*****S*****R*******

217 237
AVYTKVVHYRKWIKDTIAANP
SL***V*******

(57) Abstract

An isolated, substantially homogenous hK2 polypeptide is provided as well as isolated nucleic acid molecules encoding hK2 polypeptide, including (a) a cDNA molecule comprising the nucleotide sequence of the coding region of human hK2 gene; (b) a DNA molecule capable of hybridizing under stringent conditions to a molecule of (a); and (c) a genetic variant of any of the DNA molecules of (a) and (b) which encodes of polypeptide processing an antigenic function of naturally occurring polypeptide.

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CG	Congo	KR	Republic of Korea	SE	Sweden
CH	Switzerland	KZ	Kazakhstan	SI	Slovenia
CI	Côte d'Ivoire	LI	Liechtenstein	SK	Slovakia
CM	Cameroon	LK	Sri Lanka	SN	Senegal
CN	China	LU	Luxembourg	TD	Chad
CS	Czechoslovakia	LV	Latvia	TG	Togo
CZ	Czech Republic	MC	Monaco	TJ	Tajikistan
DE	Germany	MD	Republic of Moldova	TT	Trinidad and Tobago
DK	Denmark	MG	Madagascar	UA	Ukraine
ES	Spain	ML	Mali	US	United States of America
FI	Finland	MN	Mongolia	UZ	Uzbekistan
FR	France			VN	Viet Nam
GA	Gabon				

RECOMBINANT HK2 POLYPEPTIDE

5 Background of the Invention

The glandular kallikreins are a subgroup of serine proteases which are involved in the post-translational processing of specific polypeptide precursors to their biologically active forms. The human kallikrein gene family consists of three members: prostate-specific antigen, human glandular
 10 kallikrein, and pancreatic/renal kallikrein. See J.A. Clements, Endocr. Rev., 10, 393 (1989) and T.M. Chu et al. (U.S. Patent No. 4,446,122). A common nomenclature for these members of the tissue (glandular) kallikrein gene families was recently adopted by T. Berg et al., in Recent Progress on Kinins: Biochemistry and Molecular Biology of the Kallikrein-Kinin System. Agents
 15 and Actions Supplements, Vol. I, H. Fritz et al., eds., Birkhauser Verlag, Basel (1992), and is defined in Table I, below.

TABLE 1

20 The Human Tissue Kallikrein Gene Family (approved species designation: HSA)				
	New Designa- tion	Previous Designa- tions	mRNA/cDNA	New Protein Designation
25	hKLK1	KLK1 hRKALL	λ HK1 and phKK25 cDNAs	tissue kalli- krein (renal/ pancrease/sali- vary)
30	hKLK2	KLK2 hGK-1 hKK-3		prostate-specific glandular kalli- krein
35	hKLK3	PSA PA APS	λ HPSA-1 and PSA cDNAs	PSA (prostate- specific antigen)
40				

The DNA sequence homology between hK2 and hK3 (exon regions) is 80%, whereas the homology between hK2 and hK1 is 65%. The deduced amino acid sequence homology of hK2 to hK1 is 57%. Amino acid sequences deduced by L.J. Schedlich et al., DNA, 6, 429 (1987) and B.J. Morris, Clin. Exp. Pharmacol. Physiol. 16, 345 (1989) indicate that hK2 may be a trypsin-like serine protease, whereas hK3 (PSA) is a chymotrypsin-like serine protease. Therefore, if hK2 is indeed secretory, it may have a different physiological function than hK3.

The hK2 gene is located about 12 kbp downstream from the hK3 gene in a head-to-tail fashion on chromosome 19. (P.H. Riegman et al., FEBS Lett. 247, 123, (1989)). The similarities of gene structure and deduced amino acid sequences of these human kallikreins suggest that their evolution may involve the same ancestral gene. Most interestingly, as reported by Morris, cited supra; P. Chapdelaine, FEBS Lett. 236, 205 (1988); and Young, Biochemistry, 31, 1952 (1992), both hK2 and hK3 may be expressed only in the human prostate, while expression of hK1 is limited to the pancreas, submandibular gland, kidney, and other nonprostate tissues.

Tremendous interest has been generated in hK3 (PSA) because of the important role it plays as a marker to detect and to monitor progression of prostate carcinoma. Its usefulness as a marker is based on the elevated serum concentration of circulating hK3 proteins which are frequently associated with prostatic cancer. The serum concentration of hK3 has been found to be proportional to the cancer mass in untreated patients, but is also proportional to the volume of hyperplastic tissue in patients with benign prostatic hyperplasia (BPH). The serum levels of hK3 become reduced following prostate cancer therapy.

Despite the information which can be ascertained about hK2 from the genomic DNA sequence, very little is known about the hK2 polypeptide itself. The reason for this is that the protein has not been purified and characterized. Thus, a need exists for a method to obtain hK2 polypeptide and related polypeptides in sufficient quantity and purity for characterization and for use as therapeutic/diagnostic agents or reagents.

Summary of the Invention

The present invention provides an isolated, substantially homogenous hK2 polypeptide. As used herein, in the term "hK2 polypeptide" includes pre-pro hK2, pro hK2 and mature hK2 polypeptides. Pre-pro hK2 is
5 secreted by the cell *in vivo*, and is cleaved during secretion to yield pro hK2, which is then enzymatically cleaved in the extracellular environment to yield "mature" hK2. Most preferably, the hK2 polypeptide is contiguous in amino acid sequence with SEQ ID NO: 16, SEQ ID NO: 6 or SEQ ID NO: 10.

The present invention also provides isolated nucleic acid
10 molecules encoding hK2 polypeptide, including (a) a cDNA molecule comprising the nucleotide sequence of the coding region of the hK2 gene; (b) a DNA molecule capable of hybridizing under stringent conditions to a nucleotide sequence complementary to the nucleotide sequence of (a); and (c) a genetic variant of any of the DNA molecules of (a) and (b) which encodes
15 of polypeptide processing an antigenic function of naturally occurring hK2 polypeptide. Preferably, the nucleic acid comprises a discrete, isolated DNA or RNA molecule encoding the complete hK2 polypeptide, which can include the pre-pro, pro or mature forms. Most preferably, the nucleic acid is a DNA sequence contiguous with SEQ ID NO: 5, 7 or 9, i.e., as shown in Figs. 5, 6
20 or 7. These DNA sequences can be produced using the polymerase chain reaction (PCR), and novel oligonucleotide primers employed in the synthesis are also an embodiment of the invention.

The nucleic acid sequence also can comprise a promoter operably linked to the nucleic acid sequence. Therefore, the invention also
25 comprises a chimeric expression vector comprising the above-described nucleic acid sequence, operationally linked to control sequences recognized by a host cell transformed with the vector, as well as said transformed host cell, and methods of its preparation and use to produce recombinant hK2. Thus, the present invention also provides a method of using a nucleic acid molecule,
30 such as a cDNA clone encoding hK2 polypeptide, comprising expressing the nucleic acid molecule in a cultured host cell transformed, preferably stably transformed, with a chimeric expression vector comprising said nucleic acid

molecule operably linked to control sequences recognized by the host cell transformed with the vector, and recovering the hK2 polypeptide from the transgenic host cell, i.e., from the culture medium. As used herein, the term "chimeric" means that the vector comprises DNA from at least two different species, or comprises DNA from the same species, which is linked or
5 associated in a manner which does not occur in the "native" form of said species.

More specifically, E. coli and baculovirus insect cells systems have been employed to produce hK2 polypeptides in two forms, i.e. pre-pro
10 hK2 (pphK2) and mature hK2 (mhK2). Thus, the present invention provides the first example of the overexpression of hK2 in heterologous systems. However, although pphK2 produced in E. coli has proven to be an invaluable resource for generating antibodies to the denatured form of the protein, it is desirable to both discern the steps involved in the biosynthesis of hK2 and to
15 obtain antibodies specific for the fully processed and secreted form of the protein. Therefore, mammalian cell systems have been employed to produce hK2 polypeptides. Thus, the present invention also provides the first example of the expression of hK2 in mammalian cells and purification and characterization of the secreted protein.

20 The high degree of amino acid sequence homology of hK2 with hK3 indicates that measuring serum concentrations of both proteins may be useful in the diagnosis and monitoring of prostate cancer. For example, the antibodies developed against hK3 now used in these assays could theoretically also recognize hK2, because of mutual contamination in the antigenic
25 preparations used to develop the anti-hK3 antibodies or because of antibody cross-reactivity between these two proteins. This could account for the substantial percentage of false positive results which are observed in current hK3 assays. On the other hand, if circulating hK2 levels are also elevated above baseline levels in prostate cancer patients, detection of hK2 by hK2-
30 specific antibodies could provide an alternative, confirmatory assay for prostate cancer.

Therefore, hK2 polypeptide, as well as variants and subunits thereof, produced by the present method can be used to produce populations of antibodies that, in turn, can be used as the basis for assays to detect and quantify hK2 polypeptide (or "protein") in samples derived from tissues such as prostate carcinomas, cells such as prostate cell lines, or from fluids such as seminal fluid or blood. Thus, the present invention also provides populations of monoclonal or polyclonal antibodies that specifically bind to hK2 polypeptide, while not significantly binding to hK3. The term "significantly" is defined by reference to the comparative assays discussed below. These antibodies can also be used in affinity chromatography, to purify mammalian hK2 from natural sources. The isolated, substantially homogeneous hK2 can also be employed as a component in diagnostic assays for "native" hK2 in samples derived from human tissues or physiological fluids. For example, the recombinant hK2 can be bound to a detectable label and employed in competitive immunoassays for hK2, as described in U.S. patent application Serial No. 08/096,946, filed July 22, 1993.

As used herein with respect to the present invention, the terms "hK2 polypeptide," "hK2 protein," and "hK2" are considered to refer to identical human materials, unless otherwise indicated.

Brief Description of the Figures

Figure 1 depicts a time course study of recombinant pphK2 in Sf9 cells infected with recombinant pphK2 virus. At each of the time points cells were depleted of methionine and cysteine for 1 hour in deficient media and then supplemented with [³⁵S]-methionine and [³⁵S]-cysteine. Protein was determined by Bradford assay. Aliquots of protein (20 µg) were loaded onto a 12% Tris-Glycine SDS gel. A Phosphorimager cassette was exposed overnight. The band of interest is indicated with an arrow. w.t.: wild type.

Figure 2 depicts the detection of recombinant mhK2 in cell lysate fractions. Sf9 cells were infected either with recombinant mhK2, wild type or left uninfected for 48 hours. Methionine and cysteine pools were depleted for 1 hour in deficient media. Cells were supplemented with [³⁵S]-

methionine and [³⁵S]-cysteine for 6 hours. Cells were separated into soluble and insoluble fractions using H₂O and repeated freeze/thaw conditions. Aliquots of protein (50 µg per lane) were loaded onto a 10% Tris-Glycine SDS gel and electrophoresed. The gel was dried and exposed to x-ray film for 2 days. The band of interest is indicated with an arrow.

Figure 3 depicts the expression of recombinant pphK2 in *E. coli*. *E. coli* strain BL21 (DE3) LysS harboring pBppHK2 was grown in LB media to O.D.₆₀₀ 0.2 and incubated without (lane 2, not-induced (N)) or with (lane 3, induced (I)) 0.4 mM IPTG for 2 hrs. Cells were lysed in sample buffer and subjected to SDS/PAGE on a 4-20% gradient gel. Protein bands were visualized by staining the gel with Coomassie blue.

Figure 4 depicts the amino acid sequences of mature hK2 (deduced from cDNA sequence, SEQ ID NO: 16) and hK3 (SEQ ID NO: 1). Underlined sequences denote nonhomologous regions that can be used for preparation of antibodies specific to hK2.

Figure 5 depicts pphK2 cDNA containing a BamHI site at the 5' end and a PstI site at the 3' end (SEQ ID NO: 5) (coding strand is numbered) as well as the amino acid sequence of pre-pro hK2 encoded thereby (SEQ ID NO: 6). The amino acid sequences of pro hK2 and mature hK2 are also shown on the Figure.

Figure 6 depicts mhK2 cDNA containing an EcoRI site at the 5' end and PstI site at the 3' end (SEQ ID NO: 7), as well as the corresponding amino acid sequence (SEQ ID NO: 8) which encompasses the amino acid sequence of mhK2 polypeptide.

Figure 7 depicts pro hK2 DNA (SEQ ID NO: 9) (coding strand is numbered) and the amino acid sequence of pro hK2 (SEQ ID NO: 10).

Figure 8 depicts a gel confirming the expression of recombinant pphK2 in a mammalian cell line. AV12-pGThK2 (Lane 4-6) and AV12-pGT-d (Lane 3) clonal cell lines were grown in D10F media. About 300µl of spent medium from the above clones were concentrated and subjected to SDS/PAGE along with See Blue MW marker (lane 1) and pphK2 lysate from *E. coli* cells (lane 2). The gel was blotted onto nitrocellulose paper and

immunoblotted using a 1/1000 dilution of anti-pphK2 rabbit antiserum. HRP-goat anti-rabbit was used as the secondary probe and the blot was developed by DAB plus H_2O_2 . Lane 3 (AV12-pGT-d) is AV12 transfected with vector without insert.

5 Figure 9 depicts the DEAE chromatography of AV12 media. The sample was applied in a bicarbonate buffer, pH 8 and eluted with a salt gradient. The solid line is the A_{280} elution profile. The triangle line represents the ELISA assay of individual samples which had been dried onto microtiter plates and developed with rabbit anti-hK2 antibody.

10 Figure 10 depicts the hydrophobic interaction profile of DEAE fractions. The fractions were pooled, concentrated and applied to an HIC column in 1.2 M sodium sulfate, and eluted with a decreasing salt gradient. The solid line is A_{280} and the triangle line shows the ELISA assay profile of the fractions using rabbit anti-hK2 antibody.

15 Figure 11 depicts the Size Exclusion Chromatography of HIC purified prohK2, in particular, the A_{280} profile of 22 min peak eluted off HIC column. The 19.4 min peak appears homogeneous by SDS-PAGE. After this peak was lyophilized, the N-terminal sequence and amino acid composition confirmed its identity as the pro form of hK2.

20 Figure 12 depicts the SDS/PAGE analysis of prohK2 and PSA. 1.5 μ g of purified phK2 or PSA was boiled in sample buffer containing (R) or not containing (N) 1% BME. Samples were subjected to SDS/PAGE on a 4-20% gel. The protein bands were visualized by staining the gel with silver.

25 Detailed Description of the Invention

As used herein, the term "hK2 polypeptide" preferably encompasses the recombinant pre-pro, pro and mature hK2 polypeptides. As proposed herein, a mature hK2 polypeptide having the amino acid sequence shown in Fig. 4 (SEQ ID NO: 16), as well as "variant" polypeptides which
30 share at least 90% homology with SEQ ID NO: 16 in the regions which are substantially homologous with hK3, i.e., which regions are not identified by bars as shown in Fig. 4. Such hK2 polypeptides also possess antigenic

function in common with the mature hK2 molecule of Fig. 4, in that said polypeptides are also definable by antibodies which bind specifically thereto, but which do not cross-react with hK3 (or hK1). Preferably, said antibodies react with antigenic sites or epitopes that are also present on the mature hK2 molecule of Fig. 4. Antibodies useful to define common antigenic function are described in detail in Ser. No. 08/096,946, i.e., polyclonal antisera prepared in vivo against hK2 submit 41-56.

"Isolated hK2 nucleic acid" is RNA or DNA containing greater than 15, preferably 20 or more, sequential nucleotide bases that encode a biologically active hK2 polypeptide or a variant fragment thereof, that is complementary to the non-coding strand of the native hK2 polypeptide RNA or DNA, or hybridizes to said RNA or DNA and remains stably bound under stringent conditions. Thus, the RNA or DNA is isolated in that it is free from at least one contaminating nucleic acid with which it is normally associated in the natural source and is preferably substantially free of any other mammalian RNA or DNA. The phrase "free from at least one contaminating source nucleic acid with which it is normally associated" includes the case where the nucleic acid is reintroduced into the source or natural cell but is in a different chromosomal location or is otherwise flanked by nucleic acid sequences not normally found in the source cell. An example of isolated hK2 nucleic acid is RNA or DNA that encodes a biologically active hK2 polypeptide sharing at least 90% sequence identity with the hK3-homologous regions of the hK2 peptide of Fig. 4, as described above. The term "isolated, substantially homogenous" as used with respect to an hK2 polypeptide is defined in terms of the methodologies discussed herein below.

As used herein, the term "recombinant nucleic acid," i.e., "recombinant DNA" refers to a nucleic acid, i.e., to DNA that has been derived or isolated from any appropriate tissue source, that may be subsequently chemically altered *in vitro*, and later introduced into target host cells, such as cells derived from animal, plant, insect, yeast, fungal or bacterial sources. An example of recombinant DNA "derived" from a source, would be a DNA sequence that is identified as a useful fragment encoding

hK2, or a fragment or variant thereof, and which is then chemically synthesized in essentially pure form. An example of such DNA "isolated" from a source would be a useful DNA sequence that is excised or removed from said source by chemical means, e.g. by the use of restriction
5 endonucleases, so that it can be further manipulated, e.g., amplified, for use in the invention, by the methodology of genetic engineering.

Therefore, "recombinant DNA" includes completely synthetic DNA sequences, semi-synthetic DNA sequences, semi-synthetic DNA sequences, DNA sequences isolated from biological sources, and DNA
10 sequences derived from introduced RNA, as well as mixtures thereof. Generally, the recombinant DNA sequence is not originally resident in the genome of the host target cell which is the recipient of the DNA, or it is resident in the genome but is not expressed.

The recombinant DNA sequence, used for transformation
15 herein, may be circular or linear, double-stranded or single-stranded. Generally, the DNA sequence is in the form of chimeric DNA, such as plasmid DNA, that can also contain coding regions flanked by control sequences which promote the expression of the recombinant DNA present in the resultant cell line. For example, the recombinant DNA may itself
20 comprise a promoter that is active in mammalian cells, or may utilize a promoter already present in the genome that is the transformation target. Such promoters include the CMV promoter, as well as the SV 40 late promoter and retroviral LTRs (long terminal repeat elements). Aside from recombinant DNA sequences that serve as transcription units for hK2 or
25 portions thereof, a portion of the recombinant DNA may be untranscribed, serving a regulatory or a structural function.

"Control sequences" is defined to mean DNA sequences necessary for the expression of an operably linked coding sequence in a particular host organism. The control sequences that are suitable for
30 prokaryotic cells, for example, include a promoter, and optionally an operator sequence, and a ribosome binding site. Eukaryotic cells are known to utilize promoters, polyadenylation signals, and enhancers.

"Operably linked" is defined to mean that the nucleic acids are placed in a functional relationship with another nucleic acid sequence. For example, DNA for a presequence or secretory leader is operably linked to DNA for a polypeptide if it is expressed as a preprotein that participates in the secretion of the polypeptide; a promoter or enhancer is operably linked to a coding sequence if it affects the transcription of the sequence; or a ribosome binding site is operably linked to a coding sequence if it is positioned so as to facilitate translation. Generally, "operably linked" means that the DNA sequences being linked are contiguous and, in the case of a secretory leader, contiguous and in reading phase. However, enhancers do not have to be contiguous. Linking is accomplished by ligation at convenient restriction sites. If such sites do not exist, the synthetic oligonucleotide adaptors or linkers are used in accord with conventional practice.

Aside from recombinant DNA sequences that serve as transcription units for hK2 or portions thereof, a portion of the recombinant DNA may be untranscribed, serving a regulatory or a structural function.

The recombinant DNA to be introduced into the cells further will generally contain either a selectable marker gene or a reporter gene or both to facilitate identification and selection of transformed cells from the population of cells sought to be transformed. Alternatively, the selectable marker may be carried on a separate piece of DNA and used in a co-transformation procedure. Both selectable markers and reporter genes may be flanked with appropriate regulatory sequences to enable expression in the host cells. Useful selectable markers are well known in the art and include, for example, antibiotic and herbicide-resistance genes, such as neo, hpt, dhfr, bar, aroA, dapA and the like.

Reporter genes are used for identifying potentially transformed cells and for evaluating the functionality of regulatory sequences. Reporter genes which encode for easily assayable proteins are well known in the art. In general, a reporter gene is a gene which is not present in or expressed by the recipient organism or tissue and which encodes a protein whose expression is manifested by some easily detectable property, e.g., enzymatic activity.

Preferred genes include the chloramphenicol acetyl transferase gene (cat) from Tn9 of *E. coli*, the beta-glucuronidase gene (gus) of the *uidA* locus of *E. coli*, and the luciferase gene from firefly *Photinus pyralis*. Expression of the reporter gene is assayed at a suitable time after the DNA has been introduced into the recipient cells.

Other elements functional in the host cells, such as introns, enhancers, polyadenylation sequences and the like, may also be a part of the recombinant DNA. Such elements may or may not be necessary for the function of the DNA, but may provide improved expression of the DNA by affecting transcription, stability of the mRNA, or the like. Such elements may be included in the DNA as desired to obtain the optimal performance of the transforming DNA in the cell.

The general methods for constructing recombinant DNA which can transform target cells are well known to those skilled in the art, and the same compositions and methods of construction may be utilized to produce the DNA useful herein. For example, J. Sambrook et al., Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Laboratory Press (2d ed., 1989), provides suitable methods of construction.

The recombinant DNA can be readily introduced into the target cells by transfection with an expression vector comprising cDNA encoding hK2, for example, by the modified calcium phosphate precipitation procedure of C. Chen et al., Mol. Cell. Biol., 7, 2745 (1987). Transfection can also be accomplished by lipofectin, using commercially available kits, e.g., provided by BRL.

Suitable host cells for the expression of hK2 polypeptide are derived from multicellular organisms. Such host cells are capable of complex processing and glycosylation activities. In principle, any higher eukaryotic cell culture is workable, whether from vertebrate or invertebrate culture. Examples of invertebrate cells include plant and insect cells. Numerous baculoviral strains and variants and corresponding permissive insect host cells from hosts such as *Spodoptera frugiperda* (caterpillar), *Aedes aegypti* (mosquito), *Aedes albopictus* (mosquito), *Drosophila melanogaster* (fruitfly),

and *Bombyx mori* have been identified. See, e.g., Luckow et al., Bio/Technology, 6; 47 (1988); Miller et al., in Genetic Engineering, J. K. Setlow et al., eds., Vol. 8 (Plenum Publishing, 1986), pp. 277-279; and Maeda et al., Nature, 315:592 (1985). A variety of viral strains for transfection are
5 publicly available, e.g., the L-1 variant of *Autographa californica* NPV and the Bm-5 strain of *Bombyx mori* NPV, and such viruses may be used, preferably for transfection of *Spodoptera frugiperda* cells.

Recovery or isolation of a given fragment of DNA from a restriction digest can employ separation of the digest on polyacrylamide or
10 agarose gel by electrophoresis, identification of the fragment of interest by comparison of its mobility versus that of marker DNA fragments of known molecular weight, removal of the gel section containing the desired fragment, and separation of the gel from DNA. For example, see Lawn et al., Nucleic Acids Res., 9, 6103-6114 (1981), and Goeddel et al., Nucleic Acids Res., 8,
15 4057 (1980).

"Southern analysis" or "Southern blotting" is a method by which the presence of DNA sequences in a restriction endonuclease digest of DNA or DNA-containing composition is confirmed by hybridization to a known, labeled oligonucleotide or DNA fragment. Southern analysis typically
20 involves electrophoretic separation of DNA digests on agarose gels, denaturation of the DNA after electrophoretic separation, and transfer of the DNA to nitrocellulose, nylon, or another suitable membrane support for analysis with a radiolabeled, biotinylated, or enzyme-labeled probe as described in sections 9.37-9.52 of Sambrook et al., *supra*.

25 "Northern analysis" or "Northern blotting" is a method used to identify RNA sequences that hybridize to a known probe such as an oligonucleotide, DNA fragment, cDNA or fragment thereof, or RNA fragment. The probe is labeled with a radioisotope such as ³²P, by biotinylation or with an enzyme. The RNA to be analyzed can be usually electrophoretically
30 separated on an agarose or polyacrylamide gel, transferred to nitrocellulose, nylon, or other suitable membrane, and hybridized with the probe, using

standard techniques well known in the art such as those described in sections 7.39-7.52 of Sambrook et al., *supra*.

"Polymerase chain reaction" or "PCR" refers to a procedure or technique in which amounts of a preselected piece of nucleic acid, RNA and/or DNA, are amplified as described in U.S. Patent No. 4,683,195. Generally, sequence information from the ends of the region of interest or beyond is employed to design oligonucleotide primers. These primers will be identical or similar in sequence to opposite strands of the template to be amplified. PCR can be used to amplify specific RNA sequences, specific DNA sequences from total genomic DNA, and cDNA transcribed from total cellular RNA, bacteriophage or plasmid sequences, and the like. See generally Mullis et al., *Cold Spring Harbor Symp. Quant. Biol.*, 51, 263 (1987); Erlich, ed., *PCR Technology*, (Stockton Press, NY, 1989).

"Stringent conditions" are those that (1) employ low ionic strength and high temperature for washing, for example, 0.015 M NaCl/0.0015 M sodium citrate (SSC); 0.1% sodium lauryl sulfate (SDS) at 50°C, or (2) employ during hybridization a denaturing agent such as formamide, for example, 50% (vol/vol) formamide with 0.1% bovine serum albumin/0.1% Ficoll/0.1% polyvinylpyrrolidone/50 mM sodium phosphate buffer at pH 6.5 with 750 mM NaCl, 75 mM sodium citrate at 42°C. Another example is use of 50% formamide, 5 x SSC (0.75 M NaCl, 0.075 M sodium citrate), 50 mM sodium phosphate (pH 6.8), 0.1% sodium pyrophosphate, 5 x Denhardt's solution, sonicated salmon sperm DNA (50 µg/ml), 0.1% SDS, and 10% dextran sulfate at 42°C, with washes at 42°C in 0.2 x SSC and 0.1% SDS.

When hK2 polypeptide is expressed in a recombinant cell other than one of human origin, the hK2 polypeptide is completely free of proteins or polypeptides of human origin. However, it is necessary to purify hK2 polypeptide from recombinant cell proteins or polypeptides to obtain preparations that are substantially homogeneous as to hK2 polypeptide. For example, the culture medium or lysate can be centrifuged to remove particulate cell debris. The membrane and soluble protein fractions are then separated. The hK2 polypeptide may then be purified from the soluble

protein fraction and, if necessary, from the membrane fraction of the culture lysate. HK2 polypeptide can then be purified from contaminant soluble proteins and polypeptides by fractionation on immunoaffinity or ion-exchange columns; ethanol precipitation; reverse phase HPLC; chromatography on silica
5 or on a cation-exchange resin such as DEAE; chromatofocusing; SDS-PAGE; ammonium sulfate precipitation; gel filtration using, for example, Sephadex G-75; or ligand affinity chromatography.

Once isolated from the resulting transgenic host cells, derivatives and variants of the hK2 polypeptide can be readily prepared. For
10 example, amides of the hK2 polypeptides of the present invention may also be prepared by techniques well known in the art for converting a carboxylic acid group or precursor, to an amide. A preferred method for amide formation at the C-terminal carboxyl group is to cleave the polypeptide from a solid support with an appropriate amine, or to cleave in the presence of an alcohol,
15 yielding an ester, followed by aminolysis with the desired amine.

Salts of carboxyl groups of the hK2 polypeptide may be prepared in the usual manner by contacting the peptide with one or more equivalents of a desired base such as, for example, a metallic hydroxide base, e.g., sodium hydroxide; a metal carbonate or bicarbonate base such as, for
20 example, sodium carbonate or sodium bicarbonate; or an amine base such as, for example, triethylamine, triethanolamine, and the like.

N-acyl derivatives of an amino group of the present polypeptides may be prepared by utilizing an N-acyl protected amino acid for the final condensation, or by acylating a protected or unprotected peptide. O-
25 acyl derivatives may be prepared, for example, by acylation of a free hydroxy peptide or peptide resin. Either acylation may be carried out using standard acylating reagents such as acyl halides, anhydrides, acyl imidazoles, and the like. Both N- and O-acylation may be carried out together, if desired. In addition, the internal hK2 amino acid sequence of Fig. 4 can be modified by
30 substituting one or two conservative amino acid substitutions for the positions specified, including substitutions which utilize the D rather than L form. The invention is also directed to variant or modified forms of the hK2 polypeptide

of Fig. 4. One or more of the residues of this polypeptide can be altered, so long as antigenic function is retained. Conservative amino acid substitutions are preferred—that is, for example, aspartic-glutamic as acidic amino acids; lysine/arginine/histidine as basic amino acids; leucine/isoleucine, methionine/valine as hydrophobic amino acids; serine/glycine/alanine/threonine as hydrophilic amino acids.

Acid addition salts of the polypeptides may be prepared by contacting the polypeptide with one or more equivalents of the desired inorganic or organic acid, such as, for example, hydrochloric acid. Esters of carboxyl groups of the polypeptides may also be prepared by any of the usual methods known in the art.

Once isolated, hK2 polypeptide and its antigenically active variants, derivatives and fragments thereof can be used in assays for hK2 in samples derived from biological materials suspected of containing hK2 or anti-hK2 antibodies, as disclosed in detail in Serial No. 08/096,946. For example, the hK2 polypeptide can be labelled with a detectable label, such as via one or more radiolabelled peptidyl residues, and can be used to compete with endogenous hK2 for binding to anti-hK2 antibodies, i.e., as a "capture antigen" to bind to anti-hK2 antibodies in a sample of a physiological fluid, via various competitive immunoassay format for hK2 which uses immobilized anti-hK2 antibodies is carried out by:

- (a) providing an amount of anti-hK2 antibodies attached to a solid surface;
- (b) mixing the sample of physiological fluid to be tested with a known amount of hK2 polypeptide which comprises a detectable label, to produce a mixed sample;
- (c) contacting said antibodies on said solid surface with said mixed sample for a sufficient time to allow immunological reactions to occur between said antibodies and said hK2, and between said antibodies and said labelled polypeptide;
- (d) separating the solid surface from the mixed sample;

- (e) detecting or determining the presence or amount of labelled polypeptide either bound to the antibodies on the solid surface or remaining in the mixed sample; and
- (f) determining from the result in step (e) the presence or amount of said hK2 in said sample.

5 In another format which can detect endogenous hK2 in a sample by a competitive inhibition immunoassay, a known amount of anti-hK2 antibody is added to a sample containing an unknown amount of endogenous hK2. The known amount is selected to be less than the amount
10 required to complex all of the hK2 suspected to be present, e.g., that would be present in a sample of the same amount of physiological fluid obtained from a patient known to be prostate cancer. Next, a known amount of the hK2 polypeptide of the invention or a subunit thereof, comprising a detectable label is added. If endogenous hK2 is present in the sample, fewer antibodies
15 will be available to bind the labelled hK2 polypeptide, and it will remain free in solution. If no endogenous hK2 is present, the added labelled polypeptide will complex with the added anti-hK2 antibodies to form binary complexes. Next, the binary antibody-antigen complexes are precipitated by an anti-mammal IgG antibody (sheep, goat, mouse, etc.). The amount of radioactivity
20 or other label in the precipitate (a ternary complex) is inversely proportional to the amount of endogenous hK2 that is present in the sample, e.g., a pellet containing reduced amounts of radioactivity is indicative of the presence of endogenous hK2.

Alternatively to the conventional techniques for preparing
25 polyclonal antibodies or antisera in laboratory and farm animals, monoclonal antibodies against hK2 polypeptide can be prepared using known hybridoma cell culture techniques. In general, this method involves prepared an antibody-producing fused cell line, e.g., of primary spleen cells fused with a compatible continuous line of myeloma cells, and growing the fused cells
30 either in mass culture or in an animal species from which the myeloma cell line used was derived or is compatible. Such antibodies offer many advantages in comparison to those produced by inoculation of animals, as they

are highly specific and sensitive and relatively "pure" immunochemically. Immunologically active fragments of the present antibodies are also within the scope of the present invention, e.g., the f(ab) fragment, as are partially humanized monoclonal antibodies.

- 5 The invention will be further described by reference to the following detailed examples.

Example 1.

Construction of hK2 expression vectors

- 10 (A) Generation of recombinant baculoviruses containing pphK2 and mhK2 coding sequences

A cDNA (approximately 820 bp long) encoding the entire prepro-hK2 (pphK2) (from nucleotide #40 to #858 relative to the start site of the pphK2 transcript), as shown in Fig. 5, was synthesized from RNA of human BPH tissue using reverse-transcription polymerase chain reaction (RT-PCR) technology with a pair of hK2 specific oligonucleotide primers (5'ACGCGGATCCAGCATGTGGGACCTGGTTCTCT3' SEQ ID NO: 2 and 5'ACAGCTGCAGTTTACTAGAGGTAGGGGTGGGAC 3' SEQ ID NO:3). This cDNA was generated such that 5' and 3' ends (with respect to pphK2 sense sequence) were bracketed with BamH1 and Pst 1 sequences respectively. The cDNA was then purified by agarose gel electrophoresis, and digested with BamH1 and Pst 1 restriction enzymes. The restricted cDNA was ligated with the BamH1-Pst 1 digested pVL1393 plasmid vector and transformed into the *E. coli* HB101 strain. *E. coli* harboring pphK2 cDNA/pVL1393 plasmid vector were selected and verified by restriction enzyme mapping and DNA sequencing. Plasmid pphK2 cDNA/pVL1393 was mass-produced in *E. coli* and purified by CsCl gradient ultra-centrifugation.

cDNA encoding the mature hK2 was synthesized using PCR with the aforementioned pphK2 cDNA as the template plus a pair of hK2 oligonucleotides (5'ACGCGGATCCAGCATGTGGGACCTGGTTCTCT3' SEQ ID NO: 2 and 5'ACCGGAATTCATGATTGTGGGAGGCTGGGAGTGT3' SEQ ID NO: 4).

As noted, the 3' end oligonucleotide was the same 3' end oligonucleotide used for synthesizing the pphK2 cDNA. However, the 5' end oligonucleotide was different from the 5' oligonucleotide used for the pphK2 cDNA, and therefore generates a cDNA coding for the mature form of hK2 (mhK2), as shown in
5 Fig. 6. The mhK2 cDNA was bracketed with EcoRI and PstI sequences at the 5' and 3' ends respectively. The protein produced from the mhK2 cDNA will gain an exogenous methionine at its N-terminus. The mhK2/pVL1393 vector was generated and purified as described for pphK2/pVL1393. The DNA sequence analysis for pphK2 and mhK2 in pVL1393 showed that one
10 nucleotide (#805) has been altered (G to T) in a silent mutation.

pphK2/pVL1393 or mhK2/pVL1393 DNA (2 µg) were cotransfected with a linearized Baculogold DNA (0.5 µg; Pharmingen, San Diego, CA) into Sf9 insect cells according to Pharmingen instructions (S. Gruenwold et al., baculovirus expression vector system: Procedures and
15 Methods Manual, Pharmingen, San Diego, CA (1993)). Four to six days after the transfection, Sf9 cell spent medium containing viral particles was harvested and used to infect fresh Sf9 cells to amplify viral titers. Total RNA was isolated for Northern blot analysis of authentic pphK2 or mhK2 transcript using hK2 cDNA as a probe. Further proof of pphK2 or mhK2 transcript
20 expressed in recombinant virus infected Sf9 cells was obtained by RT-PCR and DNA sequencing. Pure recombinant baculovirus containing pphK2 or mhK2 were obtained by secondary or tertiary plaque purification protocol according to instructions from Pharmingen (S. Gruenwold et al., cited above).

25 **Example 2.**

Generation of prokaryotic expression vector

A 0.8 kb fragment representing the entire preprohK2 (pphK2) coding sequence was generated by polymerase chain reaction (PCR) using primers A (5'TATACATATGTGGGACCTGGTTCTCTCC3' SEQ ID NO.: 11)
30 and B (5'ATATGGATCCTCAGGGGTTGGCTGCGATGGT3' SEQ ID NO: 12) and plasmid pVL1393 containing pphK2 as the template. The pphK2 bacterial expression vector (pBPPHK2) was prepared by standard DNA

cloning technology, (Sambrook, cited above), to subclone this 0.8 kb fragment into the NdeI/BamHI site of the plasmid pPHS579 (a gift from Dr. H. Hsiung, Eli Lilly Co, Inc.) under the control of T7 promoter. The DNA of the entire insert plus the cloning sites was sequenced to confirm that no cloning artifacts had occurred and to ensure that no anomalies in the sequence had been generated by PCR. pBPPHK2 was transformed into E. coli BL21 (DE3)Lys S (Novagen, Inc., Madison, WI).

Example 3.

10 Generation of a mammalian expression vector

To express hK2 in mammalian cell lines, a 0.8 kb fragment representing the entire preprohK2 (pphK2) coding sequence was generated by PCR using primers
A(5'ATATGGATCCATATGTCAGCATGTGGGACCTGGTTCTCTCCA3')
15 B(5'ATATGGATCCTCAGGGGTTGGCTGCGATGGT3') (SEQ ID NO: 12)
and plasmid pVL1393 containing pphK2 as the template. The mammalian expression vector (pGThK2) was prepared using standard DNA cloning technology (Sambrook, 1989), to clone this 0.8 kb fragment into the BclI site
20 of the plasmid pGT-d (a gift from Dr. Brian Grinnell, Eli Lilly, Inc.) under control of the GBMT promoter. The DNA of the entire insert plus the cloning sites was sequenced to confirm that no cloning artifacts had occurred and to ensure that no anomalies in the sequence had been generated by PCR. AV12-664 (ATCC CRL-9595), a cell line derived from a adenovirus-induced
25 tumors in Syrian hamster, was grown in Dulbecco's modified Eagle's medium supplemented with 10% fetal bovine serum (D10F) and transfected with plasmid pGThK2 using the calcium phosphate method.

Example 4.

30 Identification of recombinant pphK2 and mhK2

A. baculovirus - insect cell system

Sf9 cells (7×10^6 /plate) were seeded onto 100 mm Corning plates with 10% fetal calf serum - Grace's medium at room temperature for 1 hr. After attachment on culture plates, cells were infected with wild type or recombinant baculovirus in serum free Excell-400 medium and incubated at 27°C. Control cells were grown in the absence of virus. At designated times (24-96 hr) cells were placed in fresh Sf9-100 media deficient of either methionine or methionine and cysteine for 45-60 min at 27°C, then incubated with Promix (0.143 mCi/plate; a mixture of [35 S]-methionine and [35 S]-cysteine; 1,000-1,400 Ci/mmol; Amersham) in serum free and methionine/cysteine deficient Sf9-100 medium (Biofluids) for 5-8 hr or 20 hr. After the end of each incubation time, cells and spent media were separated by centrifugation (1,000 rpm; Beckman J-6B; Beckman, Fullerton, CA). Cells were washed and centrifuged (13,000 rpm; Biofuge 13, Baxter) twice. The washed cells were lysed by freeze/thaw in a detergent buffer (10 mM Tris, pH 7.5; 130 mM NaCl, 1% Triton X-100, 10 mM NaF; 10 mM NaPi, 10mM Nappi, pH7.5) or H₂O and centrifuged to obtain cytosol and insoluble cellular fractions. Protein contents of the above samples were determined by either the Bradford or Lowry method (BioRad, Inc., Melville, N.Y.). The above spent media, cytosol and insoluble cellular fraction were frozen and stored separately until used. A duplicate set of samples were prepared without 35 S-labeling.

For SDS-polyacrylamide gel electrophoresis (PAGE) analysis of expression of hK2 protein in Sf9 cells, samples were added to sample buffer (U.K. Laemmli, *Nature*, 227, 680 (1970)), heated at 95°C for 5 minutes and subjected to SDS-PAGE under reducing conditions.

Northern blot analysis was routinely used to screen and isolate clonal recombinant baculoviruses expressing pphK2 or mhK2 mRNA. A comparison of the corresponding lanes in both autoradiographs of the Northern blot and photographs of ethidium bromide staining of RNA shows that mRNA for pphK2 or mhK2 was present in recombinant virus infected Sf9 but not in wild type virus-infected cells. Moreover, each of the pphK2 or mhK2 mRNA positive lanes represents RNA isolated from Sf9 cells infected

with recombinant viruses derived from a single viral plaque. Thus, the results suggest that high frequency (100%) of recombinant baculovirus containing either pphK2 or mhK2 was obtained from the above cotransfection.

Furthermore, the sequences of pphK2 or mhK2 expressed in viral infected Sf9
5 cells were confirmed by a combination of RT-PCR, cloning and DNA sequencing.

To determine whether the pphK2 protein is expressed in the insect cell Sf9, time course studies using ³⁵S-labeling of *de novo* synthesis of protein was performed and detected by SDS denaturing polyacrylamide gel
10 electrophoresis (PAGE). As seen in the autoradiograph (Fig. 1), a unique protein (about 28 KDa) was found in pphK2-recombinant virus-infected Sf9 cells at 35-74 hour post-infection. This band was missing in uninfected cells or cells infected with wild type virus. The viral polyhedron protein (about 32 KDa) was found (Fig. 1) as expected in Sf9 cells infected with wild type
15 virus, whereas it was not expressed by recombinant virus (Fig. 1). The protein was detected in cytosol when subcellular fractions (cytosol vs. insoluble fraction) was prepared by lysing cells with H₂O and freeze-thaw, whereas this 28 KD protein was detected in insoluble fraction when prepared by a detergent buffer and freeze-thaw (data not shown).

20 The mhK2 protein was also expressed in the insect cell Sf9, ³⁵S-labeling of *de novo* synthesized protein was performed. As seen in the autoradiograph (Fig. 2), a unique protein (about 28 KDa) was found in the insoluble fraction of mhK2-recombinant virus-infected Sf9 cells at 48 hours post-infection. This band was missing in uninfected cells or cells infected
25 with wild type virus. The viral polyhedron protein (about 32 KDa) was found in wild type virus-infected cells, whereas it was not expressed in cells infected with recombinant virus (Fig. 1). When the cytosol fraction was examined, no 28 KDa band was observed.

30 B. E. Coli system

Plasmid pBPPHK2 was transformed into E. coli BL21 (DE3) pLysS (Novagen, Inc., Madison, WI). This strain contains a chromosomal

copy of T7 RNA polymerase under the control of inducible LacUV5 promoter. Upon addition of IPTG (isopropyl- β -D-thiogalactopyranoside) the expression of the T7 RNA polymerase is induced which in turn activates the T7 promoter resulting in overproduction of the gene product under control of this promoter. To determine whether the product of the pphK2 gene would be expressed from pBPPHK2, single colonies of BL21 *E. Coli* transformed with pBPPHK2 were grown to O.D.₆₀₀ = 0.2 in 10 ml LB media plus ampicillin (100 μ g/ml) and induced with 0.4 mM IPTG (Sigma, Inc.). Cells were harvested 2 hours after induction by centrifugation and resuspended in 1.5 ml SDS/PAGE sample buffer (U.K. Laemmli, *Nature*, 227, 680 1970) before SDS/PAGE analysis. The cell pellet from the IPTG-induced culture was resuspended in 0.05M Tris, pH 8.0 (at 9ml/gm cell pellet) and stirred at room temperature (25°C, r.t.) for 1 hour. Lysozyme (4 mg/ml) was added to this suspension (at 1 ml/gm cell pellet) and the suspension was stirred at r.t. for 30 min followed by incubation on ice for 30 min. The suspension was sonicated for 2 min at 150 watts and centrifuged at 3000xg to isolate the inclusion bodies. Inclusion bodies were resuspended in running buffer (25 mM Tris, 192 mM glycine, 0.1% SDS) and after centrifugation both the pellet and the supernatant were analyzed by SDS/PAGE.

About 90% of the pphK2 was found to be in the supernatant fraction which indicated that pphK2 is soluble in 0.1% SDS. To prepare samples for amino acid sequence analysis, 20 μ l of inclusion body lysate was subjected to SDS/PAGE on a 4-20% gradient gel (BIO-RAD, Inc., Melville, N.Y.). The protein was blotted from the gel onto 0.2 μ PVDF paper (BIO-RAD) and stained with Coomassie blue. The protein band of interest was cut out from the blot and subjected to amino acid sequencing using a protein sequencer model 477A (Applied Biosystem, Inc., Foster City, CA).

The induced cells overproduced large amounts of a polypeptide with apparent molecular mass of about 28kd (Figure 3). Densitometric analysis indicated that this protein comprised approximately 40% of total cellular protein. The size of this protein as determined by an SDS-PAGE gel was comparable to that predicted from coding sequence for pphK2. To

confirm that this protein is pphK2, the sequence of the first 10 amino acids (MWDLVLSIAL) (SEQ ID NO: 13) from the N-terminus was determined. This sequence agrees perfectly with that deduced from the DNA sequence of pphK2 cDNA. As noted, it has different identity from the first 10 amino acids of both pphK1 (MWFLVLCLAL) (SEQ ID NO: 14) and pphK3 (MWVPVVFLTL) (SEQ ID NO: 15). It also shows that this protein is not modified or processed at the N-terminus either during or after expression in *E. coli*. These results demonstrate that we were able to accurately express pphK2 in *E. coli* from pBPPHK2.

10

C. Mammalian System

1. Isolation and Purification of protein

Plasmid pGThK2 was transformed into hamster cell line AV12-664 (ATCC-CRL-9595). To determine whether the product of the pphK2 gene would be expressed from pGThK2, AV12-pGThK2 #2 was grown in D10F + 200nM MTX. At about 60% confluency the cells were washed with Hank's balanced salt solution and resuspended in serum-free HH4 medium. The spent medium was collected after 7 days (serum-free spent medium) and stored at -20°C. Figure 8 depicts a SDS-PAGE confirming expression of recombinant pphK2 in a mammalian cell line. AV12-pGThK2 (Lane 4-6) and AV12-pGT-d (Lane 3) clonal cell lines were grown in D10F media. About 300µl of spent medium from the above clones were concentrated and subjected to SDS/PAGE along with See Blue MW marker (lane 1) and pphK2 lysate from *E. coli* cells (lane 2). The gel was blotted onto nitrocellulose paper and immunoblotted using a 1/1000 dilution of anti-pphK2 rabbit antiserum. HRP-goat anti-rabbit was used as the secondary probe and the blot was developed by DAB plus H₂O₂. Lane 3 (AV12-pGT-d) is AV12 transfected with vector without insert.

To purify the protein, the serum-free spent medium was concentrated from 5-10 fold by ultrafiltration with a 10 kDa molecular weight cutoff membrane then dialyzed overnight at 4°C versus 50 mM sodium bicarbonate, pH 8. Samples were filtered with 0.2 µ filters and then pumped

30

directly onto a TSK DEAE-5PW HPLC column (21 mm X 150 mm) at a flow rate of 5 mL/min. Buffer A contained 50 mM sodium bicarbonate, pH 7.9. Buffer B contained 50 mM sodium bicarbonate plus 0.5 M sodium chloride, pH 7.6. The elution profile shown in Figure 9 was developed with a gradient from 0-50% Buffer B over 35 min; 50-100% B from 35-40 min and isocratic elution at 100% B for 5 min before re-equilibration in Buffer A. The flow rate was 5mL/min throughout.

DEAE fractions were assayed for the presence of hK2 by ELISA using rabbit anti-pphK2 as primary antibodies. The ELISA assayed showed a peak of hK2 activity which eluted at approximately 0-2M NaCl (shown as the triangle line in Figure 9), which correlated well with the appearance of a 34 kDa band of protein seen by SDS-PAGE in the same fractions (data not shown).

Fractions with hK2 activity were pooled and concentrated by ultrafiltration with 10 kDa membranes to approximately 5-8 mL where upon solid ammonium sulfate was added to make a final concentration of 1 M. This sample was then injected onto a PolyLC. polypropyl aspartamide column, 1000A pore size, 4.6 mm X 200 mm, to resolve protein by hydrophobic interaction chromatography (HIC, see Figure 10). Buffer A was 20 mM Na phosphate, 1.2 M Na sulfate pH 6.3 and Buffer B was 50 mM Na phosphate, 5% 2-propanol, pH 7.4. The elution gradient was 0-20% B over 5 min; 20-55% B from 5-20 min, isocratic at 55% B from 20-23 min, 55-100% B from 23-25 min; isocratic at 100% B for 2 min before re-equilibration Buffer A. The flow rate was 0.7 mL/min. Greater than 90% of the A_{280} was not retained on HIC column. The main peak retained on HIC, which eluted at 22 min, also showed the highest peak of activity by ELISA assay (triangle line, Figure 10).

HIC fractions which tested positive for hK2 on ELISA were pooled, ultrafilter concentrated as above to a volume less than 1 mL then injected on a 10/30 Pharmacia S12 size exclusion column equilibrated in 100 mM ammonium acetate. The flow rate was 0.7 mL/min. When the 22 min peak from HIC was resolved by size exclusion chromatography, typically

about 80-90% of the protein A₂₈₀ eluted at 19.4 min, a retention time consistent with a protein of approximately 34 kDa (Figure 11). The only other protein peak on SEC, eluting at 16.7 min, corresponded to an about 70KDa protein seen also in previous purification steps.

5 To examine the efficiency of our purification scheme, 1.5 µg of purified pHK2 was subjected to SDS/PAGE in the presence or absence of β-mercaptoethanol (BME), and the gel was stained with silver. Results showed that the pHK2 in our sample was about 95% pure (Fig. 12). It also showed that pro-hK2 migrated at about 30 KD in the absence of BME, and it
10 migrated at about 34 kDa in the presence of BME. This pattern is similar to that observed for the PSA purified from seminal fluid (Fig. 12).

Recombinant pHK2 is recognized by rabbit anti-ppHK2, rabbit anti-PSA and a murine monoclonal antibody directed against a polypeptide covering amino acids 41-56 of hK2, when analyzed on WESTERN blots or
15 when dried down on microtiter plates. However, pHK2 was not detectable by these antibodies in sandwich assays. These results further demonstrate that the pHK2 and PSA are conformationally different and the antibodies currently available to PSA or hK2 can not detect pHK2 in its native form. Furthermore, pHK2 was not detectable by the Tandem R or free-PSA assays (immunological
20 assays for detection of PSA in serum).

A sample of the hybridoma (HK1A 523.5) secreting the murine monoclonal antibody has been deposited in the American Type Culture Collection, Rockville, MD, and assigned ATCC HB-11876.

25 2. Amino Acid Analysis and Protein Sequencing of pHK2

The peak collected off size exclusion chromatography (SEC) in ammonium acetate was lyophilized to remove the buffer then reconstituted in water. An aliquot (2.5µg) of this sample was loaded on a Porton membrane (Beckman instruments) and subjected to automated N-terminal sequence
30 analysis on an Applied Biosystems model 477A protein sequencer which yielded the following sequence:

Val-Pro-Leu-Ile-Gln-Ser-Arg-Ile-Val-Gly-Gly-Trp-Glu- (SEQ ID NO: 18). An aliquot of the same sample in water was also hydrolyzed in gaseous 6 N HCl under vacuum for 20 h at 112 °C then reconstituted in 0.1N HCl and analyzed on an Hewlett Packard Aminoquant amino acid analyzer utilizing pre-column
5 derivatization of amino acids with OPA for primary and FMOC for secondary amines.

No competing sequence was evident from the profile of amino acids released sequentially by the Edman degradation procedure. By analogy to PSA this protein is pro hK2, since the known sequence of mature PSA has
10 been shown to begin with Ileu-Val-Gly-etc and pro PSA has been postulated to have an extra 7 amino acids at the N-terminus. Amino acid analysis of this protein yielded an amino acid composition consistent with the recombinant sequence of prohK2. These results demonstrate that pphK2 was accurately expressed in the mammalian cell line AV12-664 from pGThK2.

15

Example 5.

Production of antibodies to recombinant pphK2

A. E. Coli System

To prepare pphK2 for rabbit immunization, the inclusion bodies
20 obtained from bacterial cultures of BL21 (pBpphK2) after IPTG induction as in Example 4B were resuspended in 100µl SDS/PAGE sample buffer/ml bacterial culture and electrophoresed on preparative SDS/PAGE. The pphK2 band was excised and electroeluted from the gel into running buffer (25mM Tris, 192 mM glycine, 0.1% SDS) and used as the immunogen. Two rabbits
25 were each immunized with 100µg of the immunogen in complete Freund's adjuvant and were boosted twice in three week intervals with 100µg of the immunogen in incomplete Freund's adjuvant and PBS, respectively. Rabbit anti-pphK2 sera was obtained one week following the second boost. The presence of anti-pphK2 in the rabbit antiserum was shown by ELISA (data not
30 shown). Once confirmed by this method, the highest titer antiserum was tested on Western blots using lysates from IPTG induced or non-induced cultures of BI.21 (pBpphK2). It was evident that the antiserum contained

antibodies highly specific for the pphK2 protein since a protein band at about 28kd corresponding to pphK2 was recognized only in the induced lysate. The antiserum also recognized the purified pphK2 further showing the specificity of the antibodies to pphK2. The above data demonstrate that the antibodies
5 recognize the prepro-form of hK2.

To delineate if the antiserum recognizes the mature form of hK2 (mhK2), mhK2 was expressed in *E. coli* as a glutathione S-transferase fusion protein (GST-mhK2, 58kd), and the cell lysate was immunoblotted using anit-pphK2 rabbit antiserum. It was evident that anti-pphK2 antiserum
10 recognized the GST-mhK2, demonstrating that antibodies were at least in part against the mature region of pphK2. To examine the pattern recognized in seminal fluid by anti-pphK2 antibodies, seminal fluid was prepared from pooled semen as described by Sensabaugh and Blake, *J. Urology*, 149, 1523 (1990), and immunoblotted with anti-pphK2 rabbit antiserum. The antiserum
15 recognized a major band at about 34kd plus several minor bands at lower MW. The pre-immune serum did not recognize any bands in any of the above experiments, showing that the antibodies were generated by immunization.

To determine whether there are any pphK2-specific antibodies
20 in rabbit anit-pphK2 antiserum, the antibodies cross-reacting to PSA were absorbed out of the antiserum by a PSA affinity resin. Specifically, 1ml of the sera was diluted with 1mL 100 mM HEPES, pH 7.5 and incubated with native PSA-bound Affigel-10 for 3.5 hours at 4°C. The mixture was used
25 poured into a column, the flow-through was collected and the column was washed with 30 ml HEPES buffer. Antibodies bound to the column (eluate) were eluted by acetic acid (1N, pH 4.0) and neutralized to pH 6.6 with NH₄OH. Native PSA was isolated from seminal fluid as described by Sensabaugh and Blake, cited above. ppPSA was purified from *E. coli* transformed with plasmid pPHS579 (containing ppPSA under control of T7
30 promoter) using a procedure analogous to pphK2 purification.

The flow-through and the column eluate were tested for Abs recognizing pphK2, ppPSA and native PSA (PSA isolated from seminal fluid)

using Western blot analysis. It was evident that antibodies in the untreated rabbit anti-pphK2 antiserum recognized all three proteins indicating that pphK2, ppPSA and seminal fluid-PSA share some similar epitopes. However, while the column eluate contained antibodies that recognized all three protein, the flow-through contained antibodies that recognized only pphK2. This indicates that anti-pphK2 antiserum contains pphK2-specific antibodies and these antibodies can be isolated by PSA affinity absorption. This system enabled us to generate anti-pphK2 antibodies which recognize both pphK2 and mhK2. Thus, utilizing immunogenic and pure recombinant hK2 protein, generate rabbit antiserum was generated which contains pphK2-specific antibodies, providing a valuable source for generating and screening for hK2-specific monoclonal antibodies.

These examples describe the use of three heterologous expression systems (i.e. both prokaryotic and eukaryotic) for the successful expression of the hK2 polypeptide. Thus, the method of the invention enables production of large quantities of substantially pure hK2 polypeptide. The polypeptide can be used both to study its biological functions and to produce immunodetection reagents such as labelled hK2 polypeptide, labelled fragments thereof and antibodies thereto. The immunoreagents can provide a method to purify native hK2 and to study the properties of the purified native hK2 polypeptide.

The pphK2 overproduced in *E. coli* can be readily solubilized in 0.1% SDS, thus solubility is not a problem. This is in contrast to the expression of human salivary kallikrein protein, hK1, in *E. coli*, which was found in insoluble inclusion bodies (J. Wang, et al Biochem. J., 276, 63 (1991)). In contrast, the present invention yields almost pure protein which can be purified to homogeneity by preparative SDS-PAGE. This purified recombinant pphK2 can be used for the generation of monoclonal and polyclonal antibodies.

As shown above, Baculogold viral DNA can be used to generate a recombinant baculovirus containing pphK2 or mhK2. Use of Baculogold viral DNA provides high selection of positive recombinant

baculoviruses. Indeed, Northern blot analysis showed a high frequency of recombinant virus expressing pphK2 or mhK2 mRNA. Moreover, SDS-PAGE analysis showed that both pphK2 and mhK2 recombinant viruses produced unique proteins with sizes similar to the calculated molecular weights for pphK2 or mhK2. Although the levels of the recombinant hK2 expressed in insect system may not be as high as in *E. coli*, the hK2 protein produced in baculovirus-insect system may contain the secreted form which would be more like the natural form of the protein.

Plasmids pphK2/pVL1393 in *E. coli* H13101 has been deposited in the American Type Culture Collection, Rockville, MD, USA on May 2, 1994 under the provisions of the Budapest Treaty and have been assigned accession number ATCC 69614.

The invention has been described with reference to various specific and preferred embodiments and techniques. However, it should be understood that many variations and modifications may be made while remaining within the spirit and scope of the invention.

SEQUENCE LISTING

(1) GENERAL INFORMATION:

(i) APPLICANT: Mayo Foundation for Medical Education
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Hybritech Incorporated
Tindall, Donald J.
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(ii) TITLE OF INVENTION: Recombinant HK2 Polypeptide

(iii) NUMBER OF SEQUENCES: 18

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(v) COMPUTER READABLE FORM:

(A) MEDIUM TYPE: Floppy disk
(B) COMPUTER: IBM PC compatible
(C) OPERATING SYSTEM: PC-DOS/MS-DOS
(D) SOFTWARE: PatentIn Release #1.0, Version #1.25

(vi) CURRENT APPLICATION DATA:

(A) APPLICATION NUMBER:
(B) FILING DATE:
(C) CLASSIFICATION:

(viii) ATTORNEY/AGENT INFORMATION:

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(B) REGISTRATION NUMBER: 35,561
(C) REFERENCE/DOCKET NUMBER: 150.148WO1

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(2) INFORMATION FOR SEQ ID NO:1:

- (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 237 amino acids
 (B) TYPE: amino acid
 (C) STRANDEDNESS: single
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:

```

Ile Val Gly Gly Trp Glu Cys Glu Lys His Ser Gln Pro Trp Gln Val
1      5      10
Leu Val Ala Ser Arg Gly Arg Ala Val Cys Gly Gly Val Leu Val His
20     25     30
Pro Gln Trp Val Leu Thr Ala Ala His Cys Ile Arg Asn Lys Ser Val
35     40     45
Ile Leu Leu Gly Arg His Ser Leu Phe His Pro Glu Asp Thr Gly Gln
50     55     60
Val Phe Gln Val Ser Thr Ser Phe Pro His Pro Leu Tyr Asp Met Ser
65     70     75     80
Leu Leu Lys Asn Arg Phe Leu Arg Pro Gly Asp Asp Ser Ser His Asp
85     90     95
Leu Met Leu Leu Arg Leu Ser Glu Pro Ala Glu Leu Thr Asp Ala Val
100    105    110
Lys Val Met Asp Leu Pro Thr Gln Glu Pro Ala Leu Gly Thr Thr Cys
115    120    125
Tyr Ala Ser Gly Trp Gly Ser Ile Glu Pro Glu Glu Phe Leu Thr Pro
130    135    140
Lys Lys Leu Gln Cys Val Gln Leu His Val Ile Ser Asn Asp Val Cys
145    150    155    160
Ala Gln Val His Pro Gln Lys Val Thr Lys Phe Met Leu Cys Ala Gly
165    170    175
Arg Trp Thr Gly Gly Lys Ser Thr Cys Ser Gly Asp Ser Gly Gly Pro
180    185    190
Leu Val Cys Asn Gly Val Leu Gln Gly Ile Thr Ser Trp Gly Ser Glu
195    200    205
Pro Cys Ala Leu Pro Glu Arg Pro Ser Leu Tyr Thr Lys Val Val His
210    215    220
Tyr Arg Lys Trp Ile Lys Asp Thr Ile Val Ala Asn Pro
225    230    235

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32

(2) INFORMATION FOR SEQ ID NO:2:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 32 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:2:

ACGCGGATCC AGCATGTGGG ACCTGGTTCT CT

32

(2) INFORMATION FOR SEQ ID NO:3:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 33 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:3:

ACAGCTGCAG TTTACTAGAG GTAGGGGTGG GAC

33

(2) INFORMATION FOR SEQ ID NO:4:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 34 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:4:

ACCGGAATTC ATGATTGTGG GAGGCTGGGA GTGT

34

(2) INFORMATION FOR SEQ ID NO:5:

- (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 832 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

- (ix) FEATURE:
 (A) NAME/KEY: CDS
 (B) LOCATION: 10..792

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:5:

GGATCCAGC ATG TGG GAC CTG GTT CTC TCC ATC GCC TTG TCT GTG GGG	48
Met Trp Asp Leu Val Leu Ser Ile Ala Leu Ser Val Gly	
1 5 10	
TGC ACT GGT GCC GTG CCC CTC ATC CAG TCT CGG ATT GTG GGA GGC TGG	96
Cys Thr Gly Ala Val Pro Leu Ile Gln Ser Arg Ile Val Gly Gly Trp	
15 20 25	
GAG TGT GAG AAG CAT TCC CAA CCC TGG CAG GTG GCT GTG TAC AGT CAT	144
Glu Cys Glu Lys His Ser Gln Pro Trp Gln Val Ala Val Tyr Ser His	
30 35 40 45	
GGA TGG GCA CAC TGT GGG GGT GTC CTG GTG CAC CCC CAG TGG GTG CTC	192
Gly Trp Ala His Cys Gly Gly Val Leu Val His Pro Gln Trp Val Leu	
50 55 60	
ACA GCT GCC CAT TGC CTA AAG AAG AAT AGC CAG GTC TGG CTG GGT CGG	240
Thr Ala Ala His Cys Leu Lys Lys Asn Ser Gln Val Trp Leu Gly Arg	
65 70 75	
CAC AAC CTG TTT GAG CCT GAA GAC ACA GGC CAG AGG GTC CCT GTC AGC	288
His Asn Leu Phe Glu Pro Glu Asp Thr Gly Gln Arg Val Pro Val Ser	
80 85 90	
CAC AGC TTC CCA CAC CCG CTC TAC AAT ATG AGC CTT CTG AAG CAT CAA	336
His Ser Phe Pro His Pro Leu Tyr Asn Met Ser Leu Leu Lys His Gln	
95 100 105	
AGC CTT AGA CCA GAT GAA GAC TCC AGC CAT GAC CTC ATG CTG CTC CGC	384
Ser Leu Arg Pro Asp Glu Asp Ser Ser His Asp Leu Met Leu Leu Arg	
110 115 120 125	
CTG TCA GAG CCT GCC AAG ATC ACA GAT GTT GTG AAG GTC CTG GGC CTG	432
Leu Ser Glu Pro Ala Lys Ile Thr Asp Val Val Lys Val Leu Gly Leu	
130 135 140	
CCC ACC CAG GAG CCA GCA CTG GGG ACC ACC TGC TAC GCC TCA GGC TGG	480
Pro Thr Gln Glu Pro Ala Leu Gly Thr Thr Cys Tyr Ala Ser Gly Trp	
145 150 155	
GGC AGC ATC GAA CCA GAG GAG TTC TTG CGC CCC AGG AGT CTT CAG TGT	528
Gly Ser Ile Glu Pro Glu Glu Phe Leu Arg Pro Arg Ser Leu Gln Cys	
160 165 170	

34

GTG AGC CTC CAT CTC CTG TCC AAT GAC ATG TGT GCT AGA GCT TAC TCT Val Ser Leu His Leu Leu Ser Asn Asp Met Cys Ala Arg Ala Tyr Ser 175 180 185	576
GAG AAG GTG ACA GAG TTC ATG TTG TGT GCT GGG CTC TGG ACA GGT GGT Glu Lys Val Thr Glu Phe Met Leu Cys Ala Gly Leu Trp Thr Gly Gly 190 195 200 205	624
AAA GAC ACT TGT GGG GGT GAT TCT GGG GGT CCA CTT GTC TGT AAT GGT Lys Asp Thr Cys Gly Gly Asp Ser Gly Gly Pro Leu Val Cys Asn Gly 210 215 220	672
GTG CTT CAA GGT ATC ACA TCA TGG GGC CCT GAG CCA TGT GCC CTG CCT Val Leu Gln Gly Ile Thr Ser Trp Gly Pro Glu Pro Cys Ala Leu Pro 225 230 235	720
GAA AAG CCT GCT GTG TAC ACC AAG GTG GTG CAT TAC CGG AAG TGG ATC Glu Lys Pro Ala Val Tyr Thr Lys Val Val His Tyr Arg Lys Trp Ile 240 245 250	768
AAG TAC ACC ATC GCA GCC AAC CCC TGAGTGCCCC TGTCGCCACC CTACCTCTAG Lys Tyr Thr Ile Ala Ala Asn Pro 255 260	822
TAAACTGCAG	832

(2) INFORMATION FOR SEQ ID NO:6:

- (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 261 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:6:

Met Trp Asp Leu Val Leu Ser Ile Ala Leu Ser Val Gly Cys Thr Gly
 1 5 10 15
 Ala Val Pro Leu Ile Gln Ser Arg Ile Val Gly Gly Trp Glu Cys Glu
 20 25 30
 Lys His Ser Gln Pro Trp Gln Val Ala Val Tyr Ser His Gly Trp Ala
 35 40 45
 His Cys Gly Gly Val Leu Val His Pro Gln Trp Val Leu Thr Ala Ala
 50 55 60
 His Cys Leu Lys Lys Asn Ser Gln Val Trp Leu Gly Arg His Asn Leu
 65 70 75 80
 Phe Glu Pro Glu Asp Thr Gly Gln Arg Val Pro Val Ser His Ser Phe
 85 90 95
 Pro His Pro Leu Tyr Asn Met Ser Leu Leu Lys His Gln Ser Leu Arg
 100 105 110

35

Pro Asp Glu Asp Ser Ser His Asp Leu Met Leu Leu Arg Leu Ser Glu
 115 120 125
 Pro Ala Lys Ile Thr Asp Val Val Lys Val Leu Gly Leu Pro Thr Gln
 130 135 140
 Glu Pro Ala Leu Gly Thr Thr Cys Tyr Ala Ser Gly Trp Gly Ser Ile
 145 150 155 160
 Glu Pro Glu Glu Phe Leu Arg Pro Arg Ser Leu Gln Cys Val Ser Leu
 165 170 175
 His Leu Leu Ser Asn Asp Met Cys Ala Arg Ala Tyr Ser Glu Lys Val
 180 185 190
 Thr Glu Phe Met Leu Cys Ala Gly Leu Trp Thr Gly Gly Lys Asp Thr
 195 200 205
 Cys Gly Gly Asp Ser Gly Gly Pro Leu Val Cys Asn Gly Val Leu Gln
 210 215 220
 Gly Ile Thr Ser Trp Gly Pro Glu Pro Cys Ala Leu Pro Glu Lys Pro
 225 230 235 240
 Ala Val Tyr Thr Lys Val Val His Tyr Arg Lys Trp Ile Lys Tyr Thr
 245 250 255
 Ile Ala Ala Asn Pro
 260

(2) INFORMATION FOR SEQ ID NO:7:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 760 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: double
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(ix) FEATURE:

- (A) NAME/KEY: CDS
- (B) LOCATION: 7..720

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:7:

GAATTC ATG ATT GTG GGA GGC TGG GAG TGT GAG AAG CAT TCC CAA CCC	48
Met Ile Val Gly Gly Trp Glu Cys Glu Lys His Ser Gln Pro	
1 5 10	
TGG CAG GTG GCT GTG TAC AGT CAT GGA TGG GCA CAC TGT GGG GGT GTC	96
Trp Gln Val Ala Val Tyr Ser His Gly Trp Ala His Cys Gly Gly Val	
15 20 25 30	
CTG GTG CAC CCC CAG TGG GTG CTC ACA GCT GCC CAT TGC CTA AAG AAG	144
Leu Val His Pro Gln Trp Val Leu Thr Ala Ala His Cys Leu Lys Lys	
35 40 45	

36

AAT AGC CAG GTC TGG CTG GGT CGG CAC AAC CTG TTT GAG CCT GAA GAC	192
Asn Ser Gln Val Trp Leu Gly Arg His Asn Leu Phe Glu Pro Glu Asp	
50 55 60	
ACA GGC CAG AGG GTC CCT GTC AGC CAC AGC TTC CCA CAC CCG CTC TAC	240
Thr Gly Gln Arg Val Pro Val Ser His Ser Phe Pro His Pro Leu Tyr	
65 70 75	
AAT ATG AGC CTT CTG AAG CAT CAA AGC CTT AGA CCA GAT GAA GAC TCC	288
Asn Met Ser Leu Leu Lys His Gln Ser Leu Arg Pro Asp Glu Asp Ser	
80 85 90	
AGC CAT GAC CTC ATG CTG CTC CGC CTG TCA GAG CCT GCC AAG ATC ACA	336
Ser His Asp Leu Met Leu Leu Arg Leu Ser Glu Pro Ala Lys Ile Thr	
95 100 105 110	
GAT GTT GTG AAG GTC CTG GGC CTG CCC ACC CAG GAG CCA GCA CTG GGG	384
Asp Val Val Lys Val Leu Gly Leu Pro Thr Gln Glu Pro Ala Leu Gly	
115 120 125	
ACC ACC TGC TAC GCC TCA GGC TGG GGC AGC ATC GAA CCA GAG GAG TTC	432
Thr Thr Cys Tyr Ala Ser Gly Trp Gly Ser Ile Glu Pro Glu Glu Phe	
130 135 140	
TTG CGC CCC AGG AGT CTT CAG TGT GTG AGC CTC CAT CTC CTG TCC AAT	480
Leu Arg Pro Arg Ser Leu Gln Cys Val Ser Leu His Leu Ser Asn	
145 150 155	
GAC ATG TGT GCT AGA GCT TAC TCT GAG AAG GTG ACA GAG TTC ATG TTG	528
Asp Met Cys Ala Arg Ala Tyr Ser Glu Lys Val Thr Glu Phe Met Leu	
160 165 170	
TGT GCT GGG CTC TGG ACA GGT GGT AAA GAC ACT TGT GGG GGT GAT TCT	576
Cys Ala Gly Leu Trp Thr Gly Gly Lys Asp Thr Cys Gly Gly Asp Ser	
175 180 185 190	
GGG GGT CCA CTT GTC TGT AAT GGT GTG CTT CAA GGT ATC ACA TCA TGG	624
Gly Gly Pro Leu Val Cys Asn Gly Val Leu Gln Gly Ile Thr Ser Trp	
195 200 205	
GGC CCT GAG CCA TGT GCC CTG CCT GAA AAG CCT GCT GTG TAC ACC AAG	672
Gly Pro Glu Pro Cys Ala Leu Pro Glu Lys Pro Ala Val Tyr Thr Lys	
210 215 220	
GTG GTG CAT TAC CGG AAG TGG ATC AAG TAC ACC ATC GCA GCC AAC CCC	720
Val Val His Tyr Arg Lys Trp Ile Lys Tyr Thr Ile Ala Ala Asn Pro	
225 230 235	
TGAGTGCCCC TGTCACCCC CTACCTCTAG TAAACTGCAG	760

(2) INFORMATION FOR SEQ ID NO:8:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 238 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:8:

```

Met  Ile Val Gly Gly Trp Glu Cys Glu Lys His Ser Gln Pro Trp Gln
 1      5      10      15
Val Ala Val Tyr Ser His Gly Trp Ala His Cys Gly Gly Val Leu Val
 20      25      30
His Pro Gln Trp Val Leu Thr Ala Ala His Cys Leu Lys Lys Asn Ser
 35      40      45
Gln Val Trp Leu Gly Arg His Asn Leu Phe Glu Pro Glu Asp Thr Gly
 50      55      60
Gln Arg Val Pro Val Ser His Ser Phe Pro His Pro Leu Tyr Asn Met
 65      70      75      80
Ser Leu Leu Lys His Gln Ser Leu Arg Pro Asp Glu Asp Ser Ser His
 85      90      95
Asp Leu Met Leu Leu Arg Leu Ser Glu Pro Ala Lys Ile Thr Asp Val
100      105      110
Val Lys Val Leu Gly Leu Pro Thr Gln Glu Pro Ala Leu Gly Thr Thr
115      120      125
Cys Tyr Ala Ser Gly Trp Gly Ser Ile Glu Pro Glu Glu Phe Leu Arg
130      135      140
Pro Arg Ser Leu Gln Cys Val Ser Leu His Leu Leu Ser Asn Asp Met
145      150      155      160
Cys Ala Arg Ala Tyr Ser Glu Lys Val Thr Glu Phe Met Leu Cys Ala
165      170      175
Gly Leu Trp Thr Gly Gly Lys Asp Thr Cys Gly Gly Asp Ser Gly Gly
180      185      190
Pro Leu Val Cys Asn Gly Val Leu Gln Gly Ile Thr Ser Trp Gly Pro
195      200      205
Glu Pro Cys Ala Leu Pro Glu Lys Pro Ala Val Tyr Thr Lys Val Val
210      215      220
His Tyr Arg Lys Trp Ile Lys Tyr Thr Ile Ala Ala Asn Pro
225      230      235

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(2) INFORMATION FOR SEQ ID NO:9:

- (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 766 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: double
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

- (ix) FEATURE:
 (A) NAME/KEY: CDS
 (B) LOCATION: 1..732

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:9:

GTG CCC CTC ATC CAG TCT CGG ATT GTG GGA GGC TGG GAG TGT GAG AAG	48
Val Pro Leu Ile Gln Ser Arg Ile Val Gly Gly Trp Glu Cys Glu Lys	
1 5 10 15	
CAT TCC CAA CCC TGG CAG GTG GCT GTG TAC AGT CAT GGA TGG GCA CAC	96
His Ser Gln Pro Trp Gln Val Ala Val Tyr Ser His Gly Trp Ala His	
20 25 30	
TGT GGG GGT GTC CTG GTG CAC CCC CAG TGG GTG CTC ACA GCT GCC CAT	144
Cys Gly Gly Val Leu Val His Pro Gln Trp Val Leu Thr Ala Ala His	
35 40 45	
TGC CTA AAG AAG AAT AGC CAG GTC TGG CTG GGT CGG CAC AAC CTG TTT	192
Cys Leu Lys Lys Asn Ser Gln Val Trp Leu Gly Arg His Asn Leu Phe	
50 55 60	
GAG CCT GAA GAC ACA GGC CAG AGG GTC CCT GTC AGC CAC AGC TTC CCA	240
Glu Pro Glu Asp Thr Gly Gln Arg Val Pro Val Ser His Ser Phe Pro	
65 70 75 80	
CAC CCG CTC TAC AAT ATG AGC CTT CTG AAG CAT CAA AGC CTT AGA CCA	288
His Pro Leu Tyr Asn Met Ser Leu Leu Lys His Gln Ser Leu Arg Pro	
85 90 95	
GAT GAA GAC TCC AGC CAT GAC CTC ATG CTG CTC CGC CTG TCA GAG CCT	336
Asp Glu Asp Ser Ser His Asp Leu Met Leu Leu Arg Leu Ser Glu Pro	
100 105 110	
GCC AAG ATC ACA GAT GTT GTG AAG GTC CTG GGC CTG CCC ACC CAG GAG	384
Ala Lys Ile Thr Asp Val Val Lys Val Leu Gly Leu Pro Thr Gln Glu	
115 120 125	
CCA GCA CTG GGG ACC ACC TGC TAC GCC TCA GGC TGG GGC AGC ATC GAA	432
Pro Ala Leu Gly Thr Thr Cys Tyr Ala Ser Gly Trp Gly Ser Ile Glu	
130 135 140	
CCA GAG GAG TTC TTG CGC CCC AGG AGT CTT CAG TGT GTG AGC CTC CAT	480
Pro Glu Glu Phe Leu Arg Pro Arg Ser Leu Gln Cys Val Ser Leu His	
145 150 155 160	
CTC CTG TCC AAT GAC ATG TGT GCT AGA GCT TAC TCT GAG AAG GTG ACA	528
Leu Leu Ser Asn Asp Met Cys Ala Arg Ala Tyr Ser Glu Lys Val Thr	
165 170 175	

39

GAG TTC ATG TTG TGT GCT GGG CTC TGG ACA GGT GGT AAA GAC ACT TGT	576
Glu Phe Met Leu Cys Ala Gly Leu Trp Thr Gly Gly Lys Asp Thr Cys	
180 185 190	
GGG GGT GAT TCT GGG GGT CCA CTT GTC TGT AAT GGT GTG CTT CAA GGT	624
Gly Gly Asp Ser Gly Gly Pro Leu Val Cys Asn Gly Val Leu Gln Gly	
195 200 205	
ATC ACA TCA TGG GGC CCT GAG CCA TGT GCC CTG CCT GAA AAG CCT GCT	672
Ile Thr Ser Trp Gly Pro Glu Pro Cys Ala Leu Pro Glu Lys Pro Ala	
210 215 220	
GTG TAC ACC AAG GTG GTG CAT TAC CGG AAG TGG ATC AAG TAC ACC ATC	720
Val Tyr Thr Lys Val Val His Tyr Arg Lys Trp Ile Lys Tyr Thr Ile	
225 230 235 240	
GCA GCC AAC CCC TGAGTGCCCC TGTCCCACCC CTACCTCTAG TAAA	766
Ala Ala Asn Pro	

(2) INFORMATION FOR SEQ ID NO:10:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 244 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:10:

Val Pro Leu Ile Gln Ser Arg Ile Val Gly Gly Trp Glu Cys Glu Lys	
1 5 10 15	
His Ser Gln Pro Trp Gln Val Ala Val Tyr Ser His Gly Trp Ala His	
20 25 30	
Cys Gly Gly Val Leu Val His Pro Gln Trp Val Leu Thr Ala Ala His	
35 40 45	
Cys Leu Lys Lys Asn Ser Gln Val Trp Leu Gly Arg His Asn Leu Phe	
50 55 60	
Glu Pro Glu Asp Thr Gly Gln Arg Val Pro Val Ser His Ser Phe Pro	
65 70 75 80	
His Pro Leu Tyr Asn Met Ser Leu Leu Lys His Gln Ser Leu Arg Pro	
85 90 95	
Asp Glu Asp Ser Ser His Asp Leu Met Leu Leu Arg Leu Ser Glu Pro	
100 105 110	
Ala Lys Ile Thr Asp Val Val Lys Val Leu Gly Leu Pro Thr Gln Glu	
115 120 125	
Pro Ala Leu Gly Thr Thr Cys Tyr Ala Ser Gly Trp Gly Ser Ile Glu	
130 135 140	
Pro Glu Glu Phe Leu Arg Pro Arg Ser Leu Gln Cys Val Ser Leu His	
145 150 155 160	

[illegible]

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 28 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:11:

28

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 31 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:12:

31

(2) INFORMATION FOR SEQ ID NO:13:

- (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 10 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:13:

Met Trp Asp Leu Val Leu Ser Ile Ala Leu
1 5 10

(2) INFORMATION FOR SEQ ID NO:14:

- (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 10 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:14:

Met Trp Phe Leu Val Leu Cys Leu Ala Leu
1 5 10

(2) INFORMATION FOR SEQ ID NO:15:

- (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 10 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:15:

Met Trp Val Pro Val Val Phe Leu Thr Leu
1 5 10

(2) INFORMATION FOR SEQ ID NO:16:

- (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 237 amino acids
 (B) TYPE: amino acid
 (C) STRANDEDNESS: single
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:16:

```

Ile Val Gly Gly Trp Glu Cys Glu Lys His Ser Gln Pro Trp Gln Val
1      5      10
Ala Val Trp Ser His Gly Trp Ala His Cys Gly Gly Val Leu Val His
20     25     30
Pro Gln Trp Val Leu Thr Ala Ala His Cys Leu Lys Lys Asn Ser Gln
35     40     45
Val Trp Leu Gly Arg His Asn Leu Phe Glu Pro Glu Asp Thr Gly Gln
50     55     60
Arg Val Pro Val Ser His Ser Phe Pro His Pro Leu Tyr Asn Met Ser
65     70     75     80
Leu Leu Lys His Gln Ser Leu Arg Pro Asp Glu Asp Ser Ser His Asp
85     90     95
Leu Met Leu Leu Arg Leu Ser Glu Pro Ala Lys Ile Thr Asp Val Val
100    105    110
Lys Val Leu Gly Leu Pro Thr Gln Glu Pro Ala Leu Gly Thr Thr Cys
115    120    125
Tyr Ala Ser Gly Trp Gly Ser Ile Glu Pro Glu Glu Phe Leu Arg Pro
130    135    140
Arg Ser Leu Gln Cys Val Ser Leu His Leu Leu Ser Asn Asp Met Cys
145    150    155    160
Ala Arg Ala Tyr Ser Glu Lys Val Thr Glu Phe Met Leu Cys Ala Gly
165    170    175
Leu Trp Thr Gly Gly Lys Asp Thr Cys Gly Gly Asp Ser Gly Gly Pro
180    185    190
Leu Val Cys Asn Gly Val Leu Gln Gly Ile Thr Ser Trp Gly Pro Glu
195    200    205
Pro Cys Ala Leu Pro Glu Lys Pro Ala Val Tyr Thr Lys Val Val His
210    215    220
Tyr Arg Lys Trp Ile Lys Asp Thr Ile Ala Ala Asn Pro
225    230    235

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43

(2) INFORMATION FOR SEQ ID NO:17:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 42 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:17:

ATATGGATCC ATATGTCAGC ATGTGGGACC TGGTCTCTC CA

42

(2) INFORMATION FOR SEQ ID NO:18:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 13 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:18:

Val	Pro	Leu	Ile	Gln	Ser	Arg	Ile	Val	Gly	Gly	Trp	Glu
1				5					10			

WHAT IS CLAIMED IS:

1. Isolated, substantially homogeneous pre-pro hK2 polypeptide.
- 5 2. Isolated, substantially homogenous pro hK2 polypeptide.
3. Isolated, substantially homogenous mature hK2 polypeptide.
4. An antibody that is capable of specifically binding the hK2 polypeptide
10 of claims 1, 2 or 3 and which does not bind to hK3.
5. The antibody of claim 4 which is a monoclonal antibody.
6. A hybridoma cell line producing the antibody of claim 5.
- 15 7. An isolated nucleic acid molecule encoding the polypeptides of claim 1, 2 or 3.
8. An isolated nucleic acid molecule selected from the group consisting
20 of
 - (a) cDNA comprising the nucleotide sequence of the coding region of the hK2 gene;
 - (b) DNA capable of hybridizing under stringent conditions to a nucleotide sequence complementary to the nucleotide sequence
25 of (a); and
 - (c) a genetic variant of any of the DNA of (a) and (b) which encodes a polypeptide possessing an antigenic function of naturally occurring hK2 polypeptide.
- 30 9. The nucleic acid molecule of claim 8 further comprising a promoter operably linked to the nucleic acid molecule.

10. A chimeric expression vector comprising the nucleic acid molecule of claim 7 operably linked to control sequences recognized by a host cell transformed with the vector.
- 5 11. The vector of claim 10 wherein the host cell is E. coli.
12. The vector of claim 10 wherein the host cell is a mammalian cell.
13. A host cell transformed with the vector of claim 10.
- 10 14. The host cell of claim 13 which is E. coli.
15. The host cell of claim 13 which is mammalian.
- 15 16. A method of using a nucleic acid molecule encoding a hK2 polypeptide comprising expressing the nucleic acid molecule of claim 7 in a cultured host cell stably transformed with a chimeric vector comprising said nucleic acid molecule operably linked to control sequences recognized by the host cell transformed with the vector, and
20 recovering hK2 polypeptide from the host cell.
17. The method of claim 16 wherein the host cell is E. coli.
18. The method of claim 16 wherein the host cell is mammalian.
- 25 19. The method of claim 16 wherein the nucleic acid molecule is DNA.
20. The method of claim 16 wherein the hK2 polypeptide is recovered from the host cell culture medium.

pfu/cell	35hr				52hr				74hr			
	0	0.05	0.5	w.t.	0	0.05	0.5	w.t.	0	0.05	0.5	w.t.

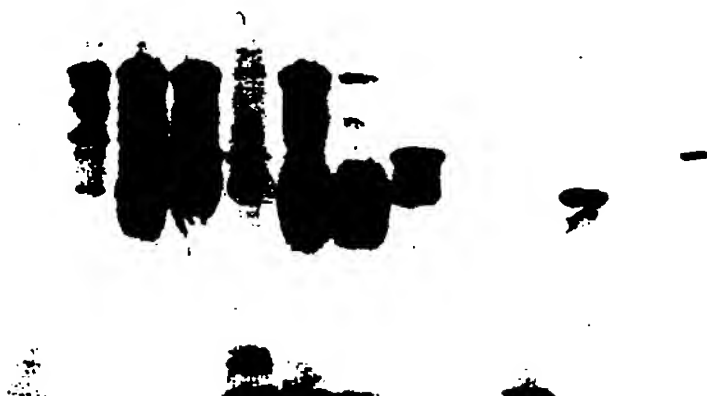


FIG. 1

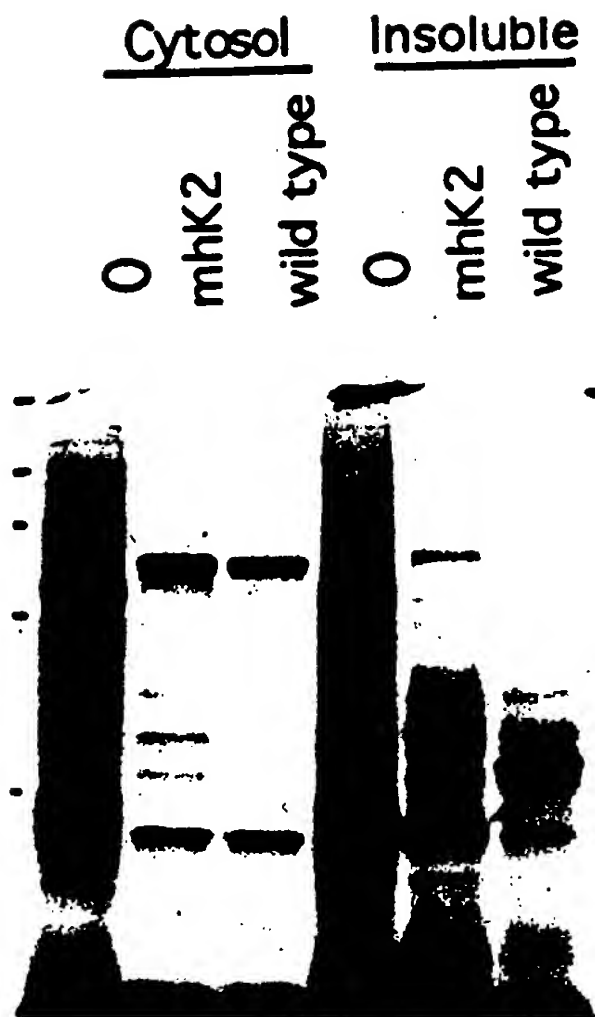


FIG. 2

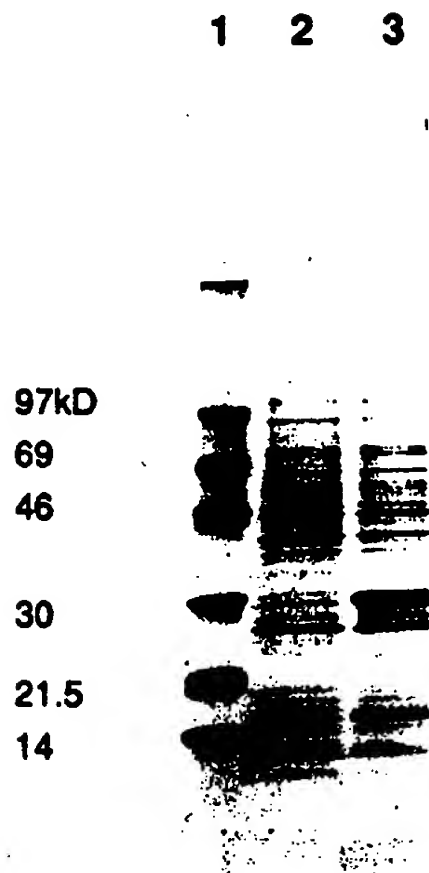


FIG. 3

hK2: IVGWECEKHSQPWQVAVVSHGWAHCGGVLVHPQWVLTAAHCLKKNQVWLGRHN
hK3: IVGWECEKHSQPWQVLA SRGRAVCGGVLVHPQWVLTAAHCIRNKS VILLGRHS

56
LFEPEDTGQRPVPSHSPHPLYNMSLLKHQSLRPDESSHDMLLRLSEPAKIT
LFHPEDTGQVFQVSTSPHPLYDMSLLKNRFLRPGDDSSHDMLLRLSEPAELT

110
153
DVVKVLGLPTQEPALGTTTCYASGWGSI EPEEFLRPRSLQCVSLHLLSNDMCA
DAVKVMDLPTQEPALGTTTCYASGWGSI EPEEFLTPKKLQCVQLHVISNDVCA

162 167
RAYSEKVTEFMLCAGLWTGGKDTCCGDSGGPLVCNGVLQGITSWGPEPCALPEKP
QVHPQKVTKFMLCAGRWTGGKSTCSGDSGGPLVCNGVLQGITSWGSEPCALPERP

217 237
AVYTKVVHYRKWKIKDTIAANP
SLYTKVVHYRKWKIKDTIVANP

FIG. 4

BamH1 ← → pphK2 → phK2
 1 GGATCCAGCATGTGGGACCTGGTTCTCTCCATCGCCTTGTCTGTGGGGTGCACTGGTGCCGTGCC
 CCTAGGTCGTACACCTGGACCAAGAGAGGTAGCGGAACAGACACCCACGTGACCACGGCACGG
 1 MetTrpAspLeuValLeuSerIleAlaLeuSerValGlyCysThrGlyAlaValPr
 → mhK2
 66 CCTCATCCAGTCTCGGATTGTGGGAGGCTGGGAGTGTGAGAAGCATTCCCAACCCCTGGCAGGTGG
 GGAGTAGGTCAGAGCCTAACACCCCTCCGACCCCTCACACTCTTCGTAAGGGTTGGGACCGTCCACC
 19 oLeuIleGlnSerArgIleValGlyGlyTrpGluCysGluLysHisSerGlnProTrpGlnValA
 131 CTGTGTACAOTCATGGATGGGCACACTGTGGGGGTGTCTGGTGCACCCCCAGTGGGTGCTCACA
 GACACATGTCAGTACCTACCCGTGTGACACCCCCACAGGACCACGTGGGGGTACCCACGAGTGT
 41 laValTyrSerHisGlyTrpAlaHisCysGlyGlyValLeuValHisProGlnTrpValLeuThr
 196 GCTGCCCATTTGCCTAAAGAAGAATAGCCAGGTCTGGCTGGGTCCGCACAACTGTTTGAGCCTGA
 CGACGGGTAACGGATTCTTCTTATCGGTCCAGACCGACCCAGCCGTGTTGGACAAACTCGGACT
 63 AlaAlaHisCysLeuLysLysAsnSerGlnValTrpLeuGlyArgHisAsnLeuPheGluProG1
 261 AGACACAGGGCCAGAGGGTCCCTGTGAGCCACAGCTTCCACACCCGCTCTACAATATGAGCCTTC
 TCTGTGTCCGGTCTCCAGGGACAGTCCGTGTGGAAGGGTGTGGGCGAGATGTTATACTCGGAAG
 84 uAspThrGlyGlnArgValProValSerHisSerPheProHisProLeuTyrAsnMetSerLeuL
 326 TGAAGCATCAAAGCCTTAGACCAGATGAAGACTCCAGCCATGACCTCATGCTGCTCCGCCTGTCA
 ACTTCGTAGTTTCGGAATCTGCTCTACTTCTGAGGTCCGTACTGGAGTACGACGAGGCGGACAGT
 106 euLysHisGlnSerLeuArgProAspGluAspSerSerHisAspLeuMetLeuLeuArgLeuSer
 391 GAGCCTGCCAAGATCACAGATGTTGTGAAGTCTCTGGGCTGCCACCCAGGAGCCAGCACTGGG
 CTCGGACGOTTCTAGTGTCTACAACACTTCCAGGACCCGGACGGGTGGGTCTCTGGTCTGACCC
 128 GluProAlaLysIleThrAspValValLysValLeuGlyLeuProThrGlnGluProAlaLeuG1
 456 GACCACCTGCTACGCCCTCAGGCTGGGGCAGCATCGAACCAGAGGAGTTCTTCCGCCCCAGGAGTC
 CTGGTGGACGATCGGAGTCCGACCCCGTCTAGCTTGGTCTCTCAAGAACCGGGGTCTCTCAG
 149 yThrThrCysTyrAlaSerGlyTrpGlySerIleGluProGluGluPheLeuArgProArgSerL
 521 TTCAGTGTGTGAGCCTCCATCTCTGTCCAATGACATGTGTCTAGAGCTTACTCTGAGAAGGTG
 AAGTCACACACTCGGAGGTAGAGGACAGGTTACTGTACACAGATCTCGAATGAGACTCTTCCAC
 171 euGlnCysValSerLeuHisLeuLeuSerAsnAspMetCysAlaArgAlaTyrSerGluLysVal
 586 ACAGAGTTTCATGTTGTGTGCTGGGCTCTGGACAGGTGOTAAAGACACTTGTGGGGGTGATTCTGG
 TGTCTCAAGTACACACAGACCGACCTGTCCACCATTCTGTGAACACCCCCACTAAGACC
 193 ThrGluPheMetLeuCysAlaGlyLeuTrpThrGlyGlyLysAspThrCysGlyGlyAspSerG1
 651 GGGTCCACTTGTCTGTATAGGTGTCTTCAAGGTATCACATCATGGGGCCCTGAGCCATGTGCCC
 CCCAGGTGAACAGACATTACCACACGAAGTTCATAGTGTAGTACCCCGGAGCTCGGTACACGGG
 214 yGlyProLeuValCysAsnGlyValLeuGlnGlyIleThrSerTrpGlyProGluProCysAlaL
 716 TGCCTGAAAAGCCTGCTGTGTACACCAAGGTGTGTCATTACCGGAAGTGGATCAAAGTACACCATC
 ACGGACTTTTCGGACGACACATGTGTTCCACCACGTAATGGCCTTCACCTAGTTTCATGTGGTAG
 236 euProGluLysProAlaValTyrThrLysValValHisTyrArgLysTrpIleLysTyrThrIle
 781 GCAGCCAACCCCTGAGTGGCCCTGTCCACCCCTACCTCTAGTAAACTGCAG
 CGTCGGTTGGGGACTCACGGGACAGCGTGGGGATGGAGATCATTTGACGTC
 258 AlaAlaAsnPro
 → Pst1

FIG. 5

1 GAATTCATGATTGTGGGAGGCTGGGAGTGTGAGAAGCATTCCCAACCC
CTTAAGTACTAACACCCTCCGACCCTCACACTCTTCGTAAGGGTTGGG
1• MetIleValGlyGlyTrpGluCysGluLysHisSerGlnPro
49 TGGCAGGTGGCTGTGTACAGTCATGGATGGGCACACTGTGGGGGTGTC
ACCGTCCACCGACACATGTCAGTACCTACCCGTGTGACACCCCCACAG
15• TrpGlnValAlaValTyrSerHisGlyTrpAlaHisCysGlyGlyVal
97 CTGGTGCACCCCCAGTGGGTGCTCACAGCTGCCCATTCCTAAAGAAG
GACCACGTGGGGGTACCCACGAGTGTGACGGGTAAACGGATTTCTTC
31• LeuValHisProGlnTrpValLeuThrAlaAlaHisCysLeuLysLys
145 AATAGCCAGGTCTGGCTGGGTGGGCACAACTGTTTGAGCCTGAAGAC
TTATCGGTCCAGACCGACCCAGCCGTGTTGGACAACTCGGACTTCTG
47• AsnSerGlnValTrpLeuGlyArgHisAsnLeuPheGluProGluAsp
193 ACAGGCCAGAGGGTCCCTGTCAGCCACAGCTTCCACACCCCGCTCTAC
TGTCCCGTCTCCAGGGACAGTCCGTGTGAAAGGTGTGGGCGAGATG
63• ThrGlyGlnArgValProValSerHisSerPheProHisProLeuTyr
241 AATATGAGCCTTCTGAAGCATCAAAGCCTTAGACAGATGAAGACTCC
TTATACCGGAAGACTTCGTAGTTTCGGAATCTGCTACTTCTGAGG
79• AsnMetSerLeuLeuLysHisGlnSerLeuArgProAspGluAspSer
289 AGCCATGACCTCATGCTGCTCCGCTGTGAGAGCCTGCCAAGATCACA
TCGGTACTGGAGTACGACGAGCGGACAGTCTCGGACGGTTCTAGTGT
95• SerHisAspLeuMetLeuLeuArgLeuSerGluProAlaLysIleThr
337 GATGTTGTGAAGGTCTGGGCTGCCACCCAGGAGCCAGCACTGGGG
CTACAACACTTCCAGGACCCGACGGGTGGGTCTCGGTCTGACCCC
111• AspValValLysValLeuGlyLeuProThrGlnGluProAlaLeuGly
385 ACCACCTGCTACGCCTCAGGCTGGGGCAGCATCGAACGAGAGGAGTTC
TGGTGGACGATGCGGAGTCCGACCCCGTCTAGCTTGGTCTCTCAAG
127• ThrThrCysTyrAlaSerGlyTrpGlySerIleGluProGluGluPhe
433 TTGCGCCCCAGGAGTCTTCAGTGTGTGAGCCTCCATCTCTGTCCAAT
AACCGCGGGTCTCAGAAGTACACACTCGGAGGTAGAGGACAGGTTA
143• LeuArgProArgSerLeuGlnCysValSerLeuHisLeuLeuSerAsn
481 GACATGTGTCTAGAGCTTACTCTGAGAAGGTGACAGAGTTCATGTTG
CTGTACACACGATCTCGAATGAGACTCTTCCACTGTCTCAAGTACAAC
159• AspMetCysAlaArgAlaTyrSerGluLysValThrGluPheMetLeu
529 TGTGCTGGGCTCTGGACAGGTGTAAGACACTTGTGGGGGTGATTCT
ACACGACCCGAGACCTGTCCACATTCTGTGAACACCCCACTAAGA
175• CysAlaGlyLeuTrpThrGlyGlyLysAspThrCysGlyGlyAspSer
577 GGGGGTCCACTTGTCTGTAATGCTGCTTCAAGGTATCACATCATGG
CCCCAGGTGAACAGACATTACCACAGAACTTCCATAGTGTAGTACC
191• GlyGlyProLeuValCysAsnGlyValLeuGlnGlyIleThrSerTrp
625 GGCCCTGAGCCATGTGCCCTGCCCTGAAAAGCCTGCTGTGTACACCAAG
CCGGGACTCGGTACACGGGACGAGCTTTTCGGACGACACATGTGGTTC
207• GlyProGluProCysAlaLeuProGluLysProAlaValTyrThrLys
673 GTGGTGCATTACCGGAAGTGGATCAAGTACACCATCGCAGCCAACCCC
CACCACGTAATGGCCTTACCTAGTTTCATGTGGTAGCGTGGTGGGG
223• ValValHisTyrArgLysTrpIleLysTyrThrIleAlaAlaAsnPro
721 TGAGTGGCCCTGTCCCACCCCTACCTCTAGTAAACTGCAG
ACTCACGGGGACAGGGTGGGGATGGAGATCATTGACGTC

FIG. 6

1 GTGCCCCATCCAGTCTCGGATTGTGGGAGGCTGGGAGTGTGAGAAGCATTCCCAACCC
CACGGGGAGTAGGTACAGACCTAACACCCCTCCGACCCCTCACACTCTTCGTAAGGGTTGGG
1> ValProLeuIleGlnSerArgIleValGlyGlyTrpGluCysGluLysHisSerGlnPro
61 TGGCAGGTGGCTGTGTACAGTCATGGATGGGCACACTGTGGGGGTGTCTGTGCACCCC
ACCGTCCACCGACACATGTCACTACCTACCCGTGTGACACCCCCACAGGACCACGTGGGG
21> TrpGlnValAlaValTyrSerHisGlyTrpAlaHisCysGlyGlyValLeuValHisPro
121 CAGTGGGTGCTCACAGCTGCCCATTTGCCTAAAGAAGAATAGCCAGGTCTGGCTGGGTTCGG
GTCACCCACGAGTGTGACGGGTAAACGGATTCTTCTTATCGGTCCAGACCACCCAGCC
41> GlnTrpValLeuThrAlaAlaHisCysLeuLysLysAsnSerGlnValTrpLeuGlyArg
181 CACAACCTGTTTGAGCCTGAAGACACAGGCCAGAGGGTCCCTGTACGCCACAGCTTCCCA
GTGTTGGACAACTCGGACTTCTGTGTCCGGTCTCCAGGGACAGTCCGTGTGAAAGGGT
61> HisAsnLeuPheGluProGluAspThrGlyGlnArgValProValSerHisSerPhePro
241 CACCCGCTCTACAATATGAGCCTTCTGAAGCATCAAAGCCTTAGACCAGATGAAGACTCC
GTGGGCGAGATGTTATACTCGGAAGACTTCGTAGTTTCGGAATCTGGTCTACTTCTGAGG
81> HisProLeuTyrAsnMetSerLeuLeuLysHisGlnSerLeuArgProAspGluAspSer
301 AGCCATGACCTCATGCTGCTCCGCTGTGAGAGCCTGCCAAGATCACAGATGTTGTGAAG
TCGGTACTGGAGTACGACGAGGCGGACAGTCTCGGACGGTCTAGTGTCTACAACACTTC
101> SerHisAspLeuMetLeuLeuArgLeuSerGluProAlaLysIleThrAspValValLys
361 GTCCTGGGCTGCCACCCAGGAGCCAGCACTGGGGACCACCTGCTACGCCCTCAGGCTGG
CAGGACCCCGGACGGGTGGGTCTCGGTCTGACCCCTGCTGGACGATCGGAGTCCGACC
121> ValLeuGlyLeuProThrGlnGluProAlaLeuGlyThrThrCysTyrAlaSerGlyTrp
421 GGCAGCATCGAACCAGAGGAGTTCTTCCGCCCCAGGAGTCTTCAGTGTGTGAGCCTCCAT
CCGTCTGTAGCTTGGTCTCTCTCAAGAACCGCGGGTCTCTCAGAAATCACACACTCGGAGGTA
141> GlySerIleGluProGluGluPheLeuArgProArgSerLeuGlnCysValSerLeuHis
481 CTCCTOTCCAATGACATGTGTCTAGAGCTTACTCTGAGAAGGTGACAGAGTTTATGTTG
GAGGACAGGTTACTGTACACAGATCTCGAATGAGACTCTTCCACTGTCTCAAGTACAAC
161> LeuLeuSerAsnAspMetCysAlaArgAlaTyrSerGluLysValThrGluPheMetLeu
541 TGTGCTGGGCTCTGGACAGGTGGTAAAGACACTTGTGGGGGTGATTCTGGGGGTCCACTT
ACACGACCCGAGACCTGTCCACCATTTCTGTGAACACCCCACTAAGACCCCAAGGTGAA
181> CysAlaGlyLeuTrpThrGlyGlyLysAspThrCysGlyGlyAspSerGlyGlyProLeu
601 GTCTGTAATGGTGTGCTTCAAGGTATCACAATCATGGGCCCCGAGCCATGTGCCCTGCCCT
CAGACATTACCACACGAAGTTCCATAGTGTAGTACCCCGGACTCGGTACACGGGACGGA
201> ValCysAsnGlyValLeuGlnGlyIleThrSerTrpGlyProGluProCysAlaLeuPro
661 GAAAAGCCTGCTGTGTACACCAAGGTGGTGCATTACCGGAAGTGGATCAAGTACACCATC
CTTTTCGACGACACATGTGTTCCACCACGTAATGGCCTTCACCTAGTTTATGTGGTAG
221> GluLysProAlaValTyrThrLysValValHisTyrArgLysTrpIleLysTyrThrIle
721 GCAGCCAACCCCTGAGTGGCCCTGTCCACCCCTACCTCTAGTAAA
CGTCGGTTGGGACTCACGGGACAGGGTGGGATGGAGATCATT
241> AlaAlaAsnPro

FIG. 7

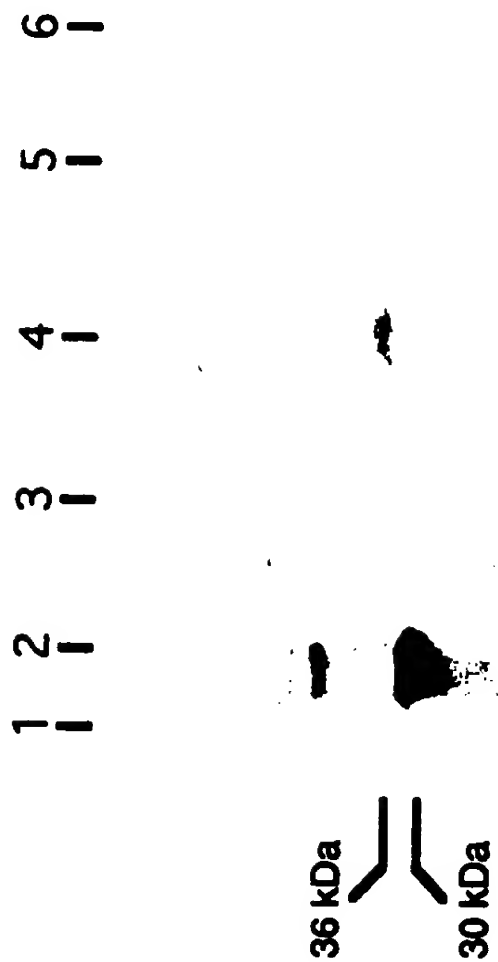


FIG. 8

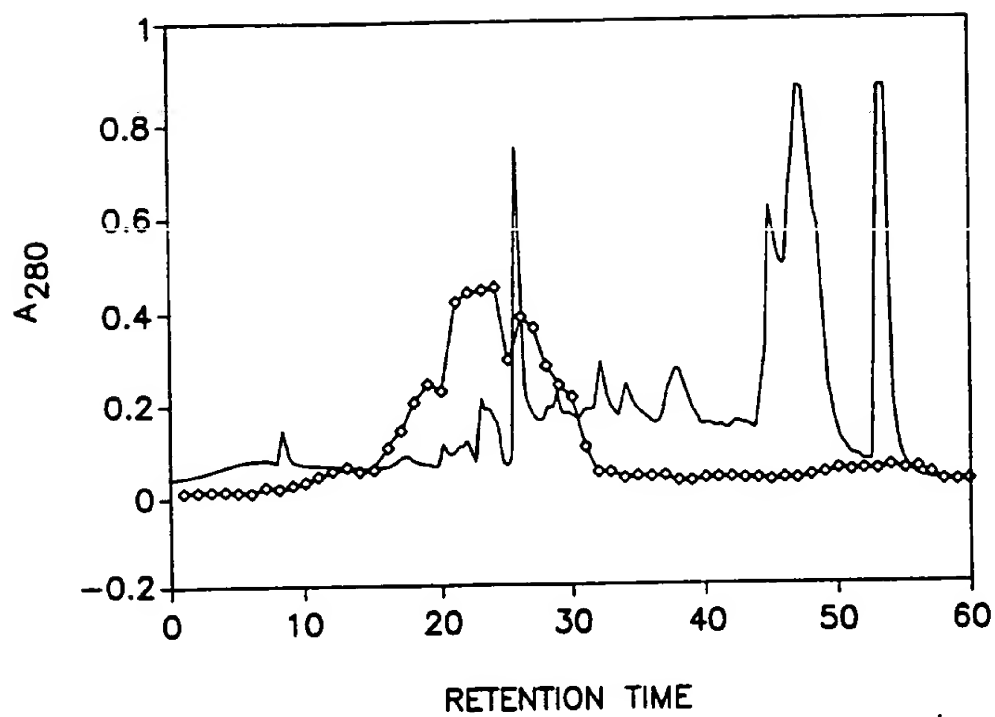


FIG. 9

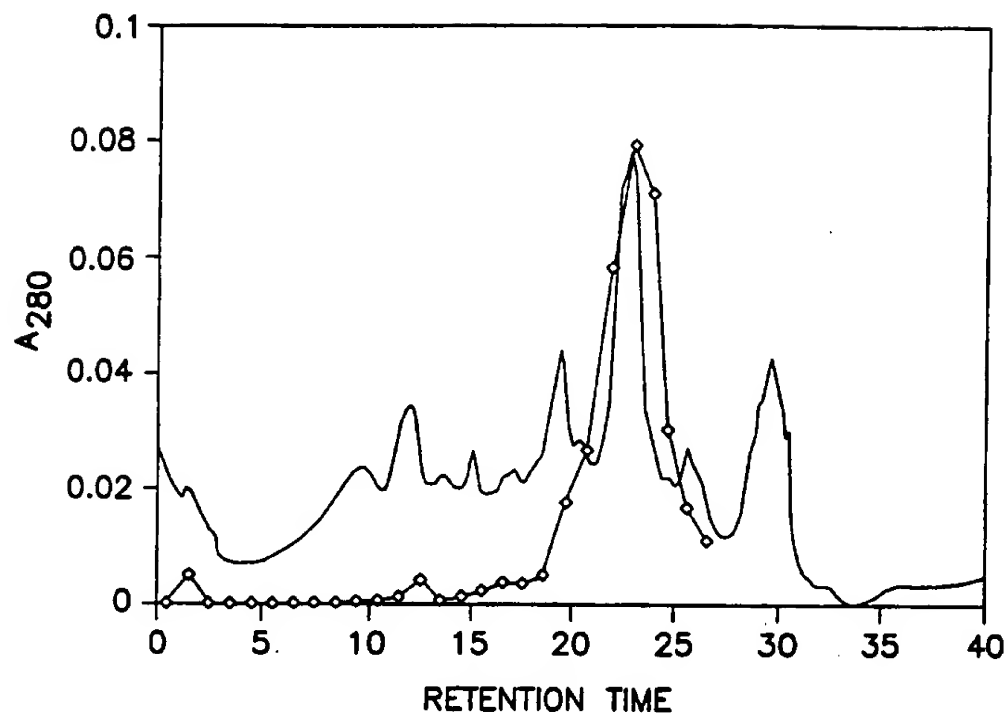


FIG. 10

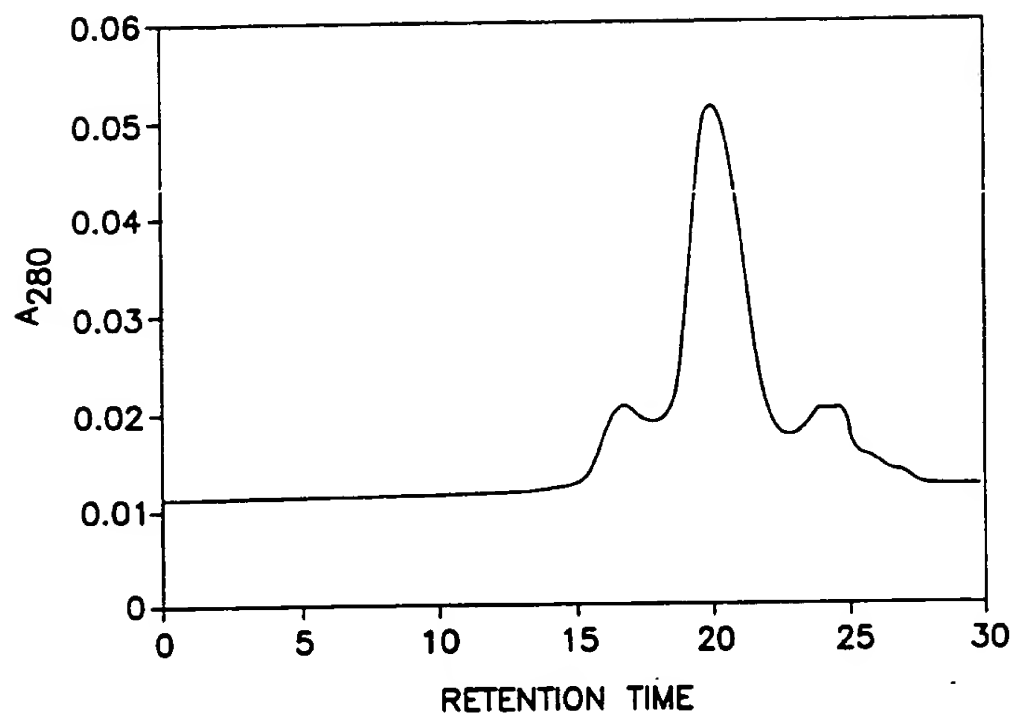


FIG. 11

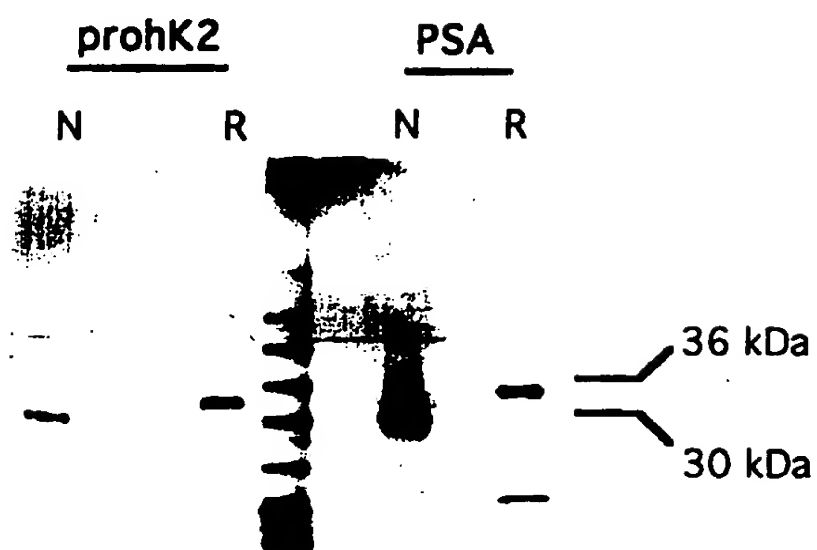


FIG. 12



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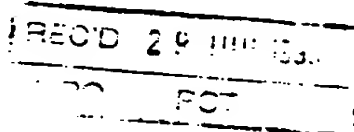
BUDAPEST TREATY ON THE INTERNATIONAL RECOGNITION OF THE DEPOSIT OF MICROORGANISMS FOR THE PURPOSES OF PATENT PROCEDURE

INTERNATIONAL FORM

RECEIPT IN THE CASE OF AN ORIGINAL DEPOSIT ISSUED PURSUANT TO RULE 7.3 AND VIABILITY STATEMENT ISSUED PURSUANT TO RULE 10.2

To: (Name and Address of Depositor or Attorney)

Hybritech, Inc.
Attn: Ru-Shya Liu
P.O. Box 289006
San Diego, CA 92196-9006



Deposited on Behalf of: Hybritech, Inc.

Identification Reference by Depositor:

ATCC Designation

Mouse hybridoma cells, HK1A523.5

HB-11878

The deposit was accompanied by: a scientific description a proposed taxonomic description
indicated above.

The deposit was received April 18, 1995 by this International Depository Authority and has been
accepted.

AT YOUR REQUEST:

☒ We will inform you of requests for the strain for 30 years.

The strain will be made available if a patent office signatory to the Budapest Treaty certifies one's right
to receive, or if a U.S. Patent is issued citing the strain and ATCC is instructed by the United States
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If the culture should die or be destroyed during the effective term of the deposit, it shall be your
responsibility to replace it with living culture of the same.

The strain will be maintained for a period of at least 30 years after the date of deposit, and for a period
of at least five years after the most recent request for a sample. The United States and many other
countries are signatory to the Budapest Treaty.

The viability of the culture cited above was tested April 24, 1995. On that date, the culture was
viable.

International Depository Authority: American Type Culture Collection, Rockville, Md. 20852 USA

Signature of person having authority to represent ATCC:


Annette L. Bada, Director, Patent Depository

Date: May 2, 1995

cc: Tim Howe



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29 JUN 1995

BUDAPEST TREATY ON THE INTERNATIONAL RECOGNITION OF THE DEPOSIT OF MICROORGANISMS FOR THE PURPOSES OF PATENT PROCEDURE

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RECEIPT IN THE CASE OF AN ORIGINAL DEPOSIT ISSUED PURSUANT TO RULE 2.1
AND VIABILITY STATEMENT ISSUED PURSUANT TO RULE 2.2

MAY 09 1994
RECEIVED

To: (Name and Address of Depositor or Attorney)

Charles Young, Ph.D.
Guggenheim 1711
Mayo Clinic
200 First Street SW
Rochester, MN 55905

Deposited on Behalf of: Charles Young, Ph.D., Mayo Clinic

Identification Reference by Depositor:

ATCC Designation

Escherichia coli HB101, pphK2/pVL1393

69614

The deposit was accompanied by: ☐ a scientific description ☒ a proposed taxonomic description indicated above.

The deposit was received May 2, 1994 by this International Depository Authority and has been accepted.

AT YOUR REQUEST:

☒ We will not inform you of requests for the strain.

The strain will be made available if a patent office signatory to the Budapest Treaty certifies one's right to receive, or if a U.S. Patent is issued citing the strain.

If the culture should die or be destroyed during the effective term of the deposit, it shall be your responsibility to replace it with living culture of the same.

The strain will be maintained for a period of at least 30 years after the date of deposit, and for a period of at least five years after the most recent request for a sample. The United States and many other countries are signatory to the Budapest Treaty.

The viability of the culture cited above was tested May 4, 1994. On that date, the culture was viable.

International Depository Authority: American Type Culture Collection, Rockville, Md. 20852 USA

Signature of person having authority to represent ATCC:

Bobbie A. Brandon Date: May 4, 1994
Bobbie A. Brandon, Head, ATCC Patent Depository

cc: Warren Woessner

Form BP4/9

INTERNATIONAL SEARCH REPORT

International Application No.

PCT/US 95/06157

A. CLASSIFICATION OF SUBJECT MATTER
 IPC 6 C12N15/57 C12N9/64 C07K16/40 C12N5/20 C12N5/10
 C12N1/21

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C12N C07K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	JOURNAL OF HYPERTENSION, vol. 6, no. s4, 1988 pages s395-s398, L. SCHEDLICH ET AL 'Kallikrein genes: cloning in man and expression in rat renal hypertension' see the whole document ---	1-20
X	MOLECULAR AND CELLULAR ENDOCRINOLOGY, vol. 76, 1991 pages 181-190, P. RIEGMAN ET AL 'Identification and androgen-regulated expression of two major human glandular kallikrein-1 (hGK-+) mRNA species' see the whole document ---	1-20

-/--

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

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T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

X document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

Y document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

A document member of the same patent family

Date of the actual completion of the international search

25 September 1995

Date of mailing of the international search report

13. 10. 95

Name and mailing address of the ISA

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 NL - 2280 HV Rijswijk
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 Fax (+ 31-70) 340-3016

Authorized officer

Van der Schaal, C

INTERNATIONAL SEARCH REPORT

Internat'l Application No

PCT/US 95/06157

C.(Communication) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	DNA, vol. 6, no. 5, 1987 pages 429-437, L. SCHEDLICH ET AL 'Primary structure of a human glandular kallikrein gene' see the whole document ---	1-20
X	FEBS, vol. 236, no. 1, August 1988 pages 205-208, P. CHAPDELAINE ET AL 'High level expression in the prostate of a human glandular kallikrein mRNA related to prostate-specific antigen' see the whole document ---	1-20
A	EP-A-0 297 913 (AMGEN INC) 4 January 1989 see the whole document ---	8-20
P,X	WO-A-95 03334 (MAYO FOUNDATION) 2 February 1995 see the whole document ---	1-20
P,X	MOL. CELL. ENDOCRINOL. (1995), 109(2), 237-41 CODEN: MCEND6; ISSN: 0303-7207, 1995 SAEDI, MOHAMMAD S. ET AL 'Overexpression of a human prostate-specific glandular kallikrein, hK2, in E. coli and generation of antibodies' see the whole document -----	1-20

INTERNATIONAL SEARCH REPORT

Int. App. No.

PCT/US 95/06157

Patent document cited in search report	Publication date	Patent (family member(s))	Publication date
EP-A-0297913	04-01-89	AT-T- 118547	15-03-95
		AU-B- 2075588	30-01-89
		DE-D- 3853023	23-03-95
		DE-T- 3853023	08-06-95
		ES-T- 2068201	16-04-95
		JP-T- 1503679	14-12-89
		PT-B- 87887	31-05-94
		WO-A- 8900192	12-01-89
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WO-A-9503334	02-02-95	AU-B- 7252594	20-02-95
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